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To Bryant 1814

for the of Anna, Bryant
6th part President

of the Medical Society
of London

A
SYLLABUS
OF
A COURSE OF
CHEMICAL LECTURES
READ AT
GUY'S HOSPITAL.

BY
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1811.

P R E F A C E.



IN every Science taught by Lectures, a Syllabus of the Course has been found of advantage to the Student. At the same time that it lays before him a comprehensive outline of the subject, and points out the several divisions and their arrangement with respect to each other, it defines the meaning and extent of scientific terms better than more verbal statements would do, and affords a convenient epitome of the rudiments of the science.

As the objects of Chemistry are various, and its views extensive, so likewise is the connection of its principles subtle, and their investigation complicated and minute: hence the aid of such a synopsis is perhaps even more requisite in this than in any other branch of Natural Philosophy. In proportion too as modern improvements have

enlarged the scope, they have increased the utility of a Prospectus to explain it; if, for instance, we take a retrospective view of the state of Chemical Science eight or ten years ago, it will appear that a number of discoveries have been made since that period, which have opened new fields of investigation, and have in some instances pointed out the imperfection of our former systems.

On these grounds it has been thought right to draw up a Syllabus of the Lectures delivered at this Hospital, and to renew it from time to time, as the progress of the Science may appear to require.

As the Course is necessarily designed for Medical Students, particular attention has been bestowed on those parts which relate to their profession. It is not, however, confined to this object alone. Chemistry is now so intimately connected with various departments of Science, and with most of the Arts and Manufactures whether useful or ornamental, that an acquaintance with it has become in some degree necessary in the general system of education; and however different the views with which the Gentleman, the Artist, and the Manufacturer may enter upon its study, each will obtain informa-

tion adapted to the particular line of his pursuit, that will amply reward him for the time he may spend in acquiring a competent knowledge of its principles.

Agreeably to this view of the matter, an endeavour has been made to point out the application of these principles, not only to the purposes of Medicine, but to most others to which Chemistry is in any way allied ; and it is hoped that, independently of advantage during the period of teaching, this Syllabus may be useful to Students when they have ceased to attend ;—that by future perusals they may not only recal such information as length of time or diversity of employment, had erased from their memories—but that it may also serve as a general outline of the science, to be filled up at their leisure—as a systematic arrangement to which they may refer whatever knowledge they shall hereafter obtain.

In preparing this Course of Lectures, considerable attention has been paid to the order and distribution of the different parts of the subject. But as Chemical Science, though making daily progress, is not yet sufficiently advanced to admit of a perfect arrangement of its parts, and as the new Nomenclature, though admirably

contrived, appears from Professor Davy's late brilliant discoveries, to have in some instances been at variance with facts, it has been deemed desirable in this Course of Lectures to guard against too strict an observance or hasty adoption of systematic views of arrangement. For this reason, the Synoptic Tables of Nomenclature, which appeared in the former Syllabus, are now suppressed; and a list of the Simple Bodies, that is, of those which, in the present state of our chemical knowledge, we are unable to decompose, has been substituted in their place.

It may be proper to mention, that although all subjects strictly belonging to Natural Philosophy are referred to the Lectures delivered at Guy's Hospital on that department of Science, yet the Chemical Lectures are occasionally illustrated by such parts of Experimental Philosophy as the subject may immediately require. By the free access, likewise, to an extensive Laboratory, the Student has an opportunity of seeing the various chemical processes conducted upon a scale corresponding with the expenditure of a large Hospital, and thereby of becoming familiarly acquainted with every step necessary in the management of such operations;—without

which the demonstrations of a Lecture-room will seldom make that impression which is necessary to fix them in his memory, and enable him to apply them with readiness and effect upon any future and distant occasion.

A Lecture on the Chemical Agencies of Electricity, and another upon Mineral Waters, are occasionally delivered during each Session. The examination of the several varieties of Urinary Calculi is also sometimes introduced. But these objects have not been treated of in the Syllabus, as they do not essentially belong to this Course of Lectures, and as their introduction is to be considered as optional.

A
SYLLABUS
OF
CHEMICAL LECTURES.

INTRODUCTION.

GENERAL division of the study of Nature, under the heads of *Natural History*, *Physics*, and *Chemistry*. Objects of these various branches of natural science generally stated. CHEMISTRY defined.

OF THE ATTRACTION OF
GRAVITATION.

General view of the subject. Cause of *Attraction* not understood. Different species or modifications of attraction. *Gravitation*.—This force is exerted between masses of matter, and often at an immense distance. Distinction be-

tween the *specific* and *absolute weight* of bodies.
Areometers. Experimental illustrations on the
 subject of specific gravity.

OF THE ATTRACTION OF COHESION OR AGGREGATION.

This attraction is exerted between the *homogeneous* particles of bodies, that is, particles of the *same kind*, and acts only in contact, or at imperceptible distances.—Its influence on the different forms of matter.—Differs from *Chemical* attraction. Distinction between the *Integrand* and *Elementary* or *Constituent* particles of bodies. Caloric acts in opposition to the attraction of aggregation.

Crystallization.—Is an effect of the attraction of aggregation. Term *Solution* explained. The formation of crystals, influenced by concentration, refrigeration, rest, atmospheric pressure, &c. and attended with various circumstances, such as change of temperature, of volume, &c. Explanation of the terms *Efflorescence* and *Deliquescence*.—Different susceptibility of crystallization in different salts—how applied to chemical analysis.

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Concise history of *Crystallography*. Sketches of *Romé de L'isle*, and *L'Abbé Haüy's* theories of the formation of crystals. *Primitive* and *secondary* forms of crystals. General idea of *truncations*, *decrements*, &c. and of the mode in which they give rise to all the varieties of form in crystallized bodies. The form of crystals, and the instrument called *Goniometer*, by means of which these forms are accurately ascertained, are important auxiliaries in mineralogical chemistry.

ON CHEMICAL ATTRACTION.

Chemical attraction or *Affinity*, like the attraction of cohesion, takes place in contact or at insensible distances, but differs from the latter in acting between *dissimilar* or *heterogeneous* particles. The result of its action is the formation of a *compound*, with properties differing from those of its constituent parts. *Chemical Analysis* consists in resolving a compound into its constituent or elementary parts. The attraction of cohesion, and chemical attraction may be generally considered as acting in opposition to each other. *Process of Solution* examined.

In most instances it is necessary, in order that chemical attraction may take place, that one of the constituents should be in the fluid state. Chemical combination produces remarkable changes in the form, colour, density, &c. of the bodies combined, and is generally attended with a change of temperature.

Bodies have amongst themselves different degrees of chemical attraction, and it is on the estimation of these different attractive forces that the Science of Chemistry rests.—Methods employed to form such estimates. *c*

Definition of the terms *Simple* and *Double elective attraction*.—Experimental illustrations.—*Precipitation*.—*Saturation*.

Berthollet's opinions on the doctrine of attraction generally stated. The quantities or masses of matter—the force of attraction of aggregation—the degree of temperature—considered as causes by which the phenomena of attraction are often modified.

Of the *limits* of chemical attraction.—Whether bodies can combine together in any proportion ; or whether they will unite only in certain determinate proportions?—Concise view of Mr. Dalton's and Dr. Wollaston's theories on the subject.

Another very common alteration is that the
medium of one of the articles is completely
altered as for Sulphur Acid & Copper
form Sulphate of Copper, & the same
Alkaline earth comes by the replacement or is
different in every respect from the original
in place.

Among the alterations of chemical alteration
there is sometimes a complete change of
form. The mixture of Lime & Sulphuric
acid in order of this to form the basis of
Lime the whole mass is converted
into a solid forming Sulphate of Lime
in place of the acid.

The cause of this is also sometimes the same
as by mixing Acids & Alkalies & when
are mixed together will produce a fluid
the action of which is to change the form of
the substance to sulphate of Lime &
by the combination of Carbonic acid gas & Ammonia
acid but we produce Carbonate of Ammonia
which is a gas forming a solid
Ammonia & the same gas will produce
Ammonia of Ammonia.

Chemical Comb is always attended with a change
of Density

By this it is seen that the Comb is quite pure the
Comb is a decomposition of water that will
be attended is a distinct combination
the Comb is a pure substance of water
water is pure. I also have seen quite
the same cases of chemical Comb there is a
change of temperature. I have seen a few
degrees of heat evolved the Comb of Chlorine
(Acid) it is seen for some it is seen greater
and instantly — added to the Comb

Black manganese from which to give a black
it is very disposed to absorb oxygen and
it also absorbs Carbonic Acid very readily
and easily converts it again

It is seen that the Comb of Chlorine is much
less a greater attraction for the Chlorine than
the Chlorine itself, it is seen being taken up
to the bottom of the cup from any Hydrogen or
Sulphate of Chlorine, it is an example of a
decomposition

There is a hole which is very deep and very large
the size of the quantity of the in. Burial of
Gada has example

There is a hole which is very deep and very large
the size of the quantity of the in. Burial of
Gada has example

CLASSIFICATION.

The principles of classification of the older chemists shown to be inconsistent with modern discoveries.—What is to be understood by the term *Element*, or *Elementary Substance*.

General view of the *simple* or *elementary bodies*. Of the principles of arrangement adopted in most modern treatises of Chemistry. Unavoidable difficulties in a systematic classification of chemical knowledge. Statement of the arrangement adopted in this course—reasons which appear to render it most eligible.

OF CALORIC.

Explanation of the terms *Heat* and *Caloric*. Various opinions on the nature of Caloric examined.—Its phenomena may be considered under two general heads, *viz.*

I.—FREE CALORIC, OR HEAT OF TEMPERATURE.

Caloric, in this state, *expands* bodies, and affects the thermometer.—*Equilibrium* of heat.—*Thermometer*.—Principle and construction of that instrument. Description of the various thermometrical scales. Nature and limits of the information

derived from these instruments.—Wedgwood's *Pyrometer*.—Peculiarity respecting the expansion of elastic fluids by heat.—Exceptions to the law of expansion by heat.*—Expansive force of freezing water.—Contraction of Argil by heat.—*Fusion and Vaporization* of bodies.—*Evaporation*, distinguished from *vaporization*.—Influence of the atmospheric pressure on vaporization.

Theory of the *Propagation and Equilibrium* of free caloric.—*Conducting powers* of bodies.—* Fluids are very slow conductors when at rest.

⑥ *Radiation* of heat.—Professor Prevost's theory of universal radiation.—*Reflexion* of heat.—Professor Pictet's apparent reflexion of cold.—Leslie's *differential thermometer*.—Bodies differ in their *reflecting, radiating, and absorbing powers*.—Explanations and experimental illustrations.—Practical applications of these principles.—*Ignition* is the ultimate effect of accumulated caloric in its free state.—Takes place in all bodies nearly at the same temperature.

II.—COMBINED CALORIC.

Caloric, when combined with bodies, may be considered in two points of view :

1st. As *Specific or Comparative heat*.—Meaning of these terms defined and illustrated by

* If a quantity of water be gradually cooled down to a certain temperature, and if that is carried still farther it will expand & if the vessel is closed it will often burst it, the force of water when it is freezing is extreme it often splits off pieces of rocks and slate, Trees &c. Clay cracks when heat is applied to this is owing to the moisture of it being taken away.

It may be said generally that all bodies, excepting Vegetable productions are fusible: When a body is rendered fluid.

The point of boiling water is 212, Alcohol 104 Hydrogen 170 by Fahrenheit's Thermometer. When water appears about to boiling present a train is heard This is owing to the agitation of the water: Then Steam arises this is vapour of water, which comes out of the vessel in a visible state, Water in high situations, as on the top of a Mountain &c, will boil at an inferior temperature when it is deprived of atmospheric pressure. If you cork up a flask of water, when the water is boiling, you take away the atmospheric pressure, then by immersing the flask of water after it has stop't boiling, into a vessel of Cold water ebullition will again commence, Glass is the best conductor of heat Copper Next, Silver is the best conductor of heat Gold &c &c Bodies differ very much in absorbing heat. Thus some things feel Cold, others warm. They are bad & good Conductors. On the appearance of frost, that when it thaws we feel it much colder This is owing to the wet air being a good conductor of heat. One to the next.

Fluids are very slow conductors of heat when at rest; this was supposed by Humphrey. They have, it is now known a very slight conducting power. This is easily seen by placing a Thermometer in a vessel of water, & pouring a thumb on its surface. Repeating it & you will not find this water become of a higher temperature & raise of Thermometer.

According to Dr Prevost's Caloric Consists of various particles. That Caloric radiates in right lines and observes of the same laws as light. The rays of Caloric can be brought into a focus. When a hot body is placed in the Centre of a Concave body the rays are converging towards that Concave body. By the same law the rays are directed in a parallel direction,



By placing a bottle of hot water in this way we can raise the column of the Thermometer very much: but by placing a body of greater heat (as a red hot bullet) we can ignite Gunpowder. After the Gunpowder has exploded you will find by applying your hand to the surface of the opposite mirror it will be very hot indeed.

Cold is capable of being reflected by a like experiment but in this the Thermometer is becoming the hot body. Various Bodies have different powers in radiating & reflecting the rays of heat. If a body was blackened it would radiate more, but reflect less. Bright & Metallic surfaces reflect most, but radiate less. Paper Surface. The reason why Metallic pots make better tea than an earthen one, is because it radiates less. Polished Metallic Substances.

Combined Caloric 1st Different bodies
have a different Capacity for Caloric
a diminished Capacity of Heat in consequence
of condensation will ignite themselves

If Ice be melted & a Thermometer be placed
in it, you will find the Thermometer remain
in the same state, this is owing to the heat
being entirely employed in changing the Ice
to a fluid state. But directly it is all melted
of course the Thermometer will rise, The diff. of
latent & Specific Caloric is that the latter is.

Latent heat requires a change of state, it undergoes
a change: hence it is called heat of fluidity or
evaporization, The Phenomena of latent heat was
found by Dr Black, he found it by filling a
vial with water, corked; boiled it, which soon
ceased boiling from the presence of the air in the
vial, the cause, was that the vapors could
not be carried
off, it was raised above the boiling point,
the cork was suddenly withdrawn, instead
of the water rushing out: it fell back to the
boiling point by the steam escaping.

3i of water in the form of steam contains more
heat than 83i of water at the boiling point (212 deg).
Alcohol of Sulphur is better than Aether for speeding
evaporation, but evaporation will go on without
the aid of these evaporating fluids. Water will
actually freeze by its own evaporation & much sooner
in a porous Vessel than being porous.

If we take a tube, with a bulb at each extremity & let that
tube be exhausted of all the air. Then wrap round one
of the bulbs with Lint soaked with Aether or Alcohol of
Sulphur, the water which is in the bulb will freeze.

experiments.—*Capacity* of bodies for caloric.—Changes of temperature produced by the condensation or rarefaction of elastic fluids.—Bodies ignited by the mere condensation of air.

2dly. In the state of *Latent heat*, also called *Heat of fluidity*.—Is that which is absorbed by bodies during their fusion or volatilization.—Account of some of Dr. Black's original experiments on the subject.—Experimental illustrations.—Cold produced by evaporation.—Cold produced by the solution of salts.—Heat evolved during the crystallization of salts.—Lavoisier's *Calorimeter*.—Application of the doctrine of latent heat to a variety of natural phenomena and artificial processes.

Of *Steam* in particular.—Distinction between *Vapour* and *Gas*.—Estimate of the elastic force of vapour.—Temperature of steam raised by increase of pressure.—*Eolipile*.—Papin's digester.—Application of steam to mechanical purposes.—Chemical history of the *Steam-Engine*.

Processes of *Distillation*—*Sublimation*—*Evaporation*—*Spontaneous evaporation*—*Hygrometer*.

Whether heat may be considered as existing in bodies in a state of *Chemical combination*, forming a third modification of combined caloric?

The extrication of heat by *friction* and *percussion*, not distinctly referable to any of the other modifications of caloric.

OF LIGHT.

Connexion between caloric and *Light*.—Nature of light.—Its *reflexion* and *refraction*.—Its *decomposition* into seven coloured rays.—These rays differ in their power of producing light.—They differ also in their power of producing heat.

Property of certain bodies to absorb light.—*Phosphorescence*.—Bolognian stone.—Canton's phosphorus.—Animal phosphorescent substances.—Luminous fish.—Glow-worm.—Lantern fly.—Rotten wood.

Connexion of *Light* with *Oxygen*.—*De-oxygenating* effects of light.—Its effects on nitrat and muriat of silver.—Mr. Wedgwood's contrivance for copying paintings on glass.—Effects of light on the animal and vegetable economy.—Rays of light, which, though not discernible, produce chemical effects.

General considerations on the effects of HEAT and LIGHT in nature.

Mercury does not freeze until it reaches 39 Degrees below Zero of Fahrenheit's Thermometer.

Soda Sulphur: 8 parts Nitrogen 5 Ammoniacal Salts 5 parts This produces an intense Cold

This produces a temperature of Zero Nitrate of Ammonia produces a great deg: of Cold when dissolved in water The Acids Diluted mixt with Snow produce a good degree of Cold a temperature of 15 below Zero. If you fill a Phial with a solution of Soda Sulphur & exclude the air from it by corking it closely, when you take out the Cork & expose the surface to the atmosphere you will see it will quickly crystallize.



OF AERIFORM SUBSTANCES OR GASSES.

These consist of a solid ponderable substance or *Base* combined in a peculiar manner with *Caloric*.

The property of the resulting gas depends upon the nature of its *Base*.

Those not condensable at common temperatures, distinguished by the name of *Permanently elastic Fluids*.

Method of ascertaining the *absolute* and *specific Gravity* of gasses, and computing their volume under different circumstances of *Pressure* and *Rarefaction*.

Elasticity of the air.—Theory of the *Air-Pump*.

Of Oxygen Gas.

Called formerly *Dephlogisticated*, *Pure*, *Empyreal*, or *Vital Air*.

Somewhat heavier than *Common Air*.—100 cubical inches at the temp. of 60 *Farenh.* and at 30 inches *Bar. press.* weigh about 33,8 grains.

Forms a constituent part of the *Atmosphere*—Serves the purposes of *Respiration* and *Combustion* in an eminent degree—Has a powerful affinity with a great variety of substances, particularly inflammable bodies, and when the new compounds are rapidly formed, *Caloric* and *Light* are frequently elicited.

Theory of *Combustion*—*Pyrophorus*.

To *Sulphur*, *Phosphorus*, and *Charcoal*, in the act of burning, it communicates *acid* properties ; which are more or less distinct, as the combustion or decomposition of the Gas has been more or less complete.

The basis, therefore, of this fluid was formerly considered as the universally *Acidifying Principle*, and hence denominated *Oxygen* :—hence also the terms *Oxyd*, *Oxydation*, *Oxygenation*, &c.

The base of *Oxygen Gas* has a strong attraction for *Caloric*, but this is overcome or modified by the attraction of the particles of some solid substances.

It destroys the splendour and tenacity of the *Metallic Bodies* when they are exposed to it at a high temperature, and gives them an increase of weight proportioned to its own consumption : hence the modern theory of *Calcination*.



But when Henry no longer found it in his power
to give the day in front of him in the
dark he was forced to turn in the morning - putting
a cork lightly in at last. He took it out
in the old way - but he was not

Oxygen combines in diff. proportions

but does not combine Oxygen these things.

If Iron is added it attracts Oxygen most readily
and so it is that the hydrogen gas is the most
— property of Hydrogen.

Calcium, for instance, has a great attraction for oxygen —
and they combine their bodies in long tubes that expand to 7
times.

If the hydrogen, methane are heated with Chlorine and
then exposed to the light.

If Hydrochloric acid is heated in a sealed, common air
vial first over a flame, then Oxygen and if you
immerse it in water and you will have a solution.

In process of decomposition, important to the action of the bodies
and the effect it produces in the
transformation — he thought that he had not a chance of
this in a gasometer — He quickly moved on this point
any experiments he found that an animal could only live
certain time in a certain quantity of air — and that
an animal must be used in which he could have been
out of for a long time without fresh breathing apparatus.
Could he make up that he wanted the mechanism of many bodies
the supply — he found that the gas in air was 1/5 of
oxygen. He then by making it by various means by the
and will not support life, as he found it would go on too
rapidly. He finally found that the difference of color in
arteries & veins was due to the action of the oxygen
in the blood — he found that the blood in the arteries
was oxygenated and in the veins it was deoxygenated. He
found that the blood in the arteries was red and in the veins
it was black. The Oxygen was not in the mouth of

the same is very distinctly

for every last inch of Oxygen gas

When examining the air found they found that it was very like
that of Carbonic acid

Now also all I see the Oxygen is not that same
but same also the quantity of Azote remaining

During the course of Oxygen lost in the process is
replaced with an equal volume of Carbonic acid

Now the same experiment is a further proof
that a mixture of Hydrogen & Oxygen is equal to the amount
of heat as the Hydrogen

During this process the absorption of Carbonic
acid is less than it is when in air mixture

Now also the quantity of Oxygen is taken out the
air and with an equal amount of its Carbonic acid

the same is very distinctly

but that is also because the heat is very far
from being taken out of the air

The decomposition of Oxygen is greater in heat
than in the atmosphere

Now the Oxygen is not taken out of the air
but the quantity of Carbonic acid is off in 24 hours

the same is very distinctly a further proof
that the heat is not taken out of the air
but the heat is taken out of the air

Protoxyd—the first degree of *Oxygenation*.

Peroxyd—the highest degree.

Importance of *Oxygen Gas* in the process of *Respiration*—Analogy between this process and *Combustion*. — *Carbonic Acid* the result ; —for every cubic inch of *Carbonic Acid* formed, there is a loss of a cubic inch of *Oxygen* in the air respired.—Produces a remarkable change in *Blood*: hence the supposed function of the *Lungs*, and origin of *Animal Heat*.

Its purity best ascertained by exposure to a solution of green *Sulphate* or *Muriate* of *Iron* saturated with *Nitrous Gas*, or by mixture with the *Gas* itself: *Eudiometer*.

Obtained most readily from *Metallic Oxyds*, and compounds of the *Nitric* and other acids by the application of heat, and in a state of great purity from *Oxymuriate* of *Potash*.

Of Azote or Nitrogen Gas.

Called formerly *Phlogisticated Air*, and *Mofete*.

Remains after the abstraction of *Oxygen Gas* from *Atmospherical Air*, by exposure of the latter to *Sulphuret* of *Potash*, to a mixture of *Iron Filings* and *Sulphur*, &c.

Obtained also by mixing *Oxymuriatic Acid* with *Ammoniacal Gas*.

When pure has a faint smell, but no taste, and is rather lighter than *Atmospherical Air*.

100 cubical inches, *Therm.* 60° *Bar.* 30° weigh about 30, 4 grs.

If inspired is instantaneously destructive of *Animal Life*, and extinguishes *Flame*; but does not impede *Vegetation*.

Differs from most other *Gases* in being but in a small degree capable of absorption by *Water*, or by the liquid forms either of *Acids* or *Alkalies*.

Evolved in the decomposition of *Animal* and *Vegetable Substances*, of which it forms a constituent part.

When mixed in a certain proportion with *Oxygen Gas* and exposed to the *Electrical Spark*, it produces *Nitric Acid*; and by union with *Hydrogen Gas* it forms *Ammonia*—from the former of these properties it takes the name of *Nitrogen*.

Of the means thought to be employed by nature to maintain a due proportion between this and *Oxygen Gas* in the composition of the *Atmosphere*.

Dr. William Briggs

The statement of Petrus is incorporated in the original.
sent to the Secretary in 1895 -

The 1st of the year of 1881 was quiet & calm. The
 weather was fine. The weather was fine. The weather was fine.
 The weather was fine. The weather was fine. The weather was fine.
 The weather was fine. The weather was fine. The weather was fine.

3. Express your own Court much better. It is
to be noted in a Court of Justice
and is usually made by it. The end of the
law is to be left to it.

at 212 Water is attracted by heat takes the form of
Steam

Water can only be got pure by distilling it is a glass
retort & then very slowly almost be able to take
the drops in
Water contains soft substances

Water in a slight degree evaporates

Recomposes glass of made into a state of vapor
and applied to it

Hydrogen has a strong attraction for lime &
combines with some of the particles by the action of
lime

Of Hydrogen Gas, or Inflammable Air.

Found in a disengaged and impure state in *Coal Mines*, on the surface of *Stagnant Waters*, and rising through the waters of certain *Springs*.

Obtained also artificially from *animal, vegetable, and bituminous* matter by distillation; from *Essential Oils, Alcohol, Æthers, &c.* by the application of heat; and from *Ammoniacal Gas* by means of the *Electric Spark*.

But purest from the decomposition of *Water* by *Metals*, as above stated; or during their solution in *diluted Acids*.

When pure, between 11 and 12 times lighter than *Atmospherical Air*: hence the construction of *Aërostatic Machines*.

Smell adventitious.

100 cubical inches at a Temp: 60 *Farenht.* and 30 inches *Bar. press.* weigh 2. 8 grs.

Combined with *Oxygen* in the proportion of about 15 *Hydrogen* to 85 *Oxygen* by weight, it forms water, from the decomposition of which these fluids may be both obtained in a gaseous state, and therefore considered as its elements.

Hence the effects of *Water* in promoting *Combustion*, and in the *Oxydation* of *Metals, &c. &c.*

In the combustion of *Hydrogen* with *Oxygen Gas*, the purity of the resulting water depends on the slowness or rapidity with which the process is conducted.

On inspiration proves noxious to *Animal life*, apparently by the exclusion of the *respirable* part of the *Atmosphere*.

In its nascent state it is found to be capable of combining both with *Oxygen* and *Azote*, with the one producing *Water* as in the case of combustion, and with the other *Ammonia*.

Carburetted Hydrogen Gas—formerly called, *Heavy Inflammable Air*—procured by passing *Hydrogen Gas* over red hot *Charcoal*—afforded by subjecting *vegetables* and *Inflammable Substances* to a strong heat—different kinds of *Carburetted Hydrogen*—*Olefiant Gas*—*Gas Lights*.

Sulphuretted Hydrogen Gas—Methods of procuring—Combines with *Alkaline Bases*—Is absorbed by water—The solution precipitates some metals and not others.

Phosphuretted Hydrogen Gas—One of the most combustible substances known—Absorbable by water—Methods of procuring.

Hydrogen Gas is capable of holding *Arsenic* in solution.

It may be decomposed by a common Electrical
machine Take a Tube filled with water. Cork it at
the ends & put a small gold wire thro' it. Letting the
wire communicate with an Electrical machine
an spark will come from its strong attraction
oxygen the water will become muddy...
It may be completely decomposed by making
Steam pass thro' a red hot Tube... in which
there is piece of iron wire which attracts
Oxygen & the Hydrogen passes over & may be
burned.

The inconvenience arising from Gas lights in a room
for it is not only unpleasant but unwholesome -
Hydrogen will take up Sulphur for many Sulphur
retted Hydrogen gas To prevent this pour a small
quantity of diluted Muriatic Acid on a small
portion of Sulphurated Potash. It is Sulphurated
Hydrogen which is so offensive in common Stoves
ditches &c. it is very Volatile. The bad smell of Stoves
&c. Mr Allen says may be abated by Lime & Lime
Phosphorus combined with Hyd: forming Phosphor-
retted Hydrogen Gas it is made by putting a
piece of Phosphorus in Hydrogen gas directly
this Gas is exposed to Oxygen it becomes
a compound of Phosphorus & Lime when put
in the water it will take fire

Hydrogen can hold a portion of Ammonia in
it, it is produced by heating
an Zinc



Nitrous Oxid Gas can't be respired longer than
four Minutes without losing entirely
voluntary Motion. Some persons after
taking it are remarkably grave & sullen. Some
are very lively and in fact almost deranged

Inflammable Substances require a higher
temperature to burn in this Gas than in
Hydrogen. A Piece of Sulphur must be
heated some time before it will burn at
the Nitrous Oxid Gas.

If peroxide this Gas takes Nitrate of Ammonia
which consists of Ammonia (composed of Hydrogen
and Azote) Nitric Acid (composed of Azote & Oxygen)
the result will be by distilling the Salt we
shall have ..

Phosphorus will readily decompose this Gas
and burn most splendidly.

Of Nitrous Oxyd Gas.

Procured from the decomposition of *Nitrate of Ammonia* with a gentle heat. Consists of *Oxygen* and *Azote* in intimate union. In some of its properties resembles *Acids*.

100 cubical inches at a middle temperature and pressure, weigh about 50 grains—Is decomposed by combustible substances at a very high temperature—Soluble in double its volume of *Water*, to which it communicates a sweetish taste.

Most inflammable substances require a high temperature to burn in it—Is not acted upon by *Nitrous Gas*.

Remarkable for the intoxicating effects which it produces in respiration.



OF SALTS.

These characterized by being *sapid* or of a *saline taste* and readily *soluble* in water ; further remarkable for their disposition to act on, or to be acted upon by most other substances, as also for the regularity of the forms which they are in most instances found capable of assuming—*Crystallization*.

Their solution in water for the most part accompanied by a diminution of temperature ; hence the usual means of producing *artificial Cold*.

On exposure to air generally either receive or impart moisture : hence said to be *deliquescent* or *efflorescent*.

When heated, the greater number undergo either *watery* or *igneous Fusion*—some are volatilized, and many more or less completely decomposed.

They are divided into *simple* and *compounded*—the more simple are *Alkalies* and *Acids*—the compounded, such as result from the union of these with each other, *Neutral Salts*,—or of the acids with earths and metallic oxyds, *Earthy* and *Metallic Salts*.

In some Salts there is a great portion of Water
In Alum there is a very large portion of Water
In Sal Rochelle there is also a large portion
of water

The particles of a Saline body having a disposition
to attract each other they enable us to
purify this Salt

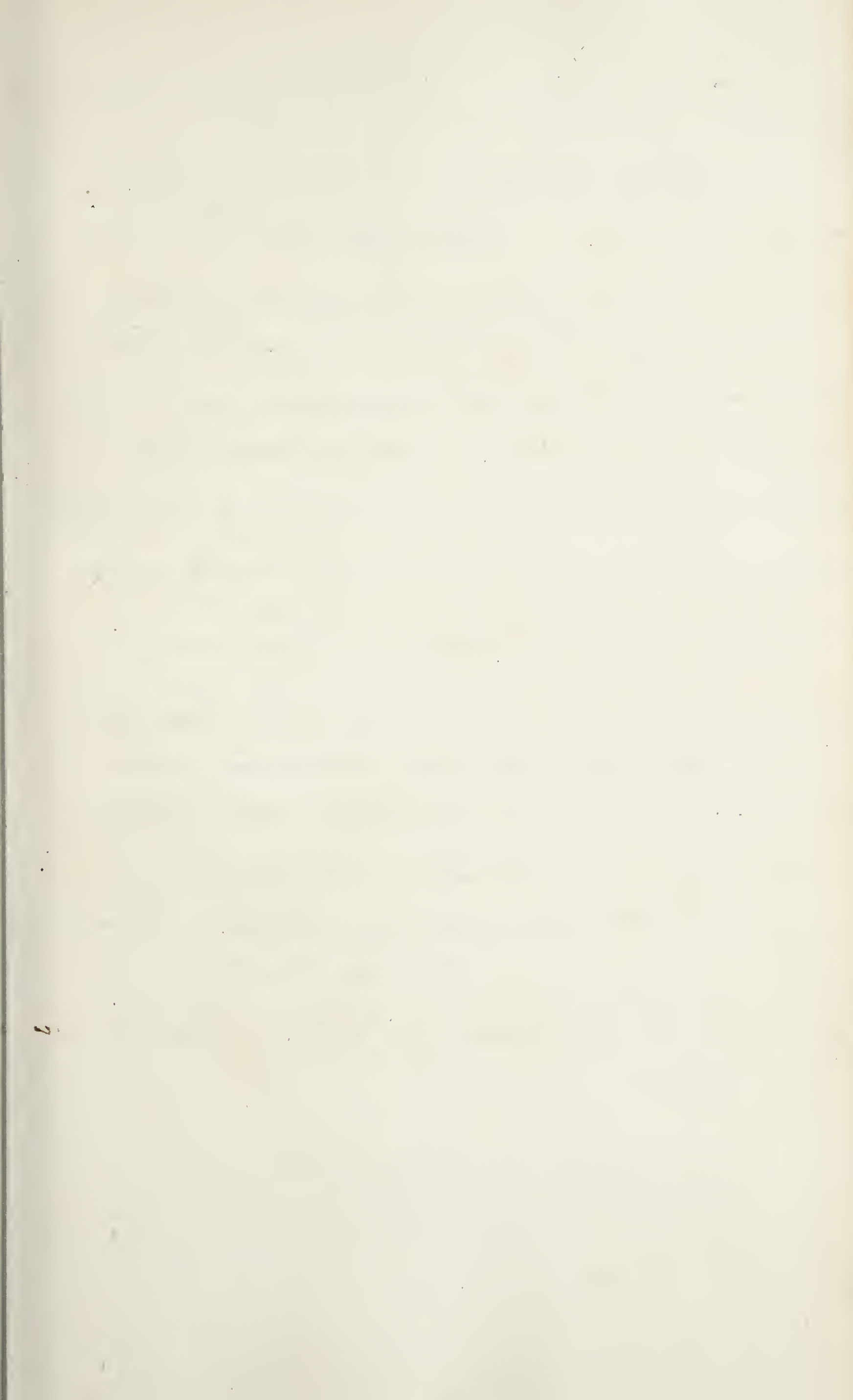
If we have a Solution of a Saline substance all
we have to do is to apply heat to evaporate the water
when on suffering it to cool Crystals will form
If you evaporate too much of the water the Crystals
will be irregular The appearance of a Pellicle is
a sign that Crystals will form when the tempera-
ture of the Solution is down to the tempera-
ture of the laboratory you will know that the
process of Crystallization is finished In
the Vessel it is situated place pieces of Stick
across the Vessel for the Crystals to form
on If the Crystals are colored you have
only to dissolve them Crystals again in water
and evaporate it again when you will obtain
pure Crystals Crystallization also affords
an opportunity of separating different Salts
which are not readily soluble in water by evaporat-
ing it cautiously you will easily separate
them - The most soluble will crystallize first

We can make Salts Crystallize directly
on applying some substance which has a
strong Attraction for water. Take a Solution
of Sulphate of Potash & pour a little Alcohol into it
when a precipitation of Crystals will take place.

Some Salts some Salts will alter the
Crystallized form, by being exposed to the Air.
The Sulphate of Potash will effloresce, Caustic
Potash will deliquesce.

Some Salts may be sublimed. Some Salts are decomposed
by heat: Saline Substances form a grand Division
Simple & Compound

We name earthy Salts, & Metallic Salts.



We know an Alkaline solution take a piece
of Paper which has been dyed in Turmeric ..
it will change its color. Syrup of Violet will
change it green - The different ^{alkalis} have no
action on each other. They have a strong
attraction for Moisture. A good test for
Alkalies is a bit of Litmus paper redened by
Vinegar if it be dip't in that state into an
Alkaline solution it will recover its blue color
Alkalies in their free state combine
with Oils & form soap

The alkalis have a Peculiar effect on Animal
Matter therefore they are called Caustics
When combined with Acids this power
is destroyed. The term till lately was
confined to these 3 substances Potash,
Soda & ammonia. but it is now
found to be in many other Substances.

OF ALKALIES.

Distinguished by a pungent, and lixivial or urinous *Taste*, and by changing most vegetable *blue* colours to *green*, and many of the vegetable *yellow* colours to *brown*.

Have a strong attraction for *moisture*.

Produce no alteration on each other, but manifest an extensive disposition to unite with other bodies. With *Acids* they constitute *Neutral Salts*.

They precipitate from their solutions most of the *Earths*, and all the *Metallic Oxyds*, several of which they are capable of re-dissolving.

With *unctuous substances* they form *Soaps*; with *Silex* and *Alumine* various kinds of *Glass* and *Porcelain*; with *Sulphur*, *Alkaline Hepars* or *Sulphurets*.

The substances most perfectly *Alkaline* are *Potash*, *Soda*, and *Ammonia*.

The two former, till lately considered as simple bodies, but discovered by Professor Davy to consist of peculiar *Metallic Bases* combined with *Oxygen*.

Ammonia remarkable for its volatility in a moderate temperature—a compound of *Hydrogen* and *Azote*.

Many of the *Earths* also have distinct alkaline properties, more especially *Barytes*, *Strontian* and *Lime*.

Of Potash.

Form, *concrete*. Taste, *extremely pungent*, *Caustic*. Colour, *white*. When perfectly dry, is a non-conductor of *Electricity*. In this state consists, according to the experiments of Professor Davy, of a peculiar metal *Potassium*—combined with *Oxygen* and a portion of *Water* intimately combined.

Potassium is considerably lighter than *Water*—rapidly absorbs *Oxygen*—is capable of two states of oxidation, *Protoxyd* and *Peroxyd*—exists in the latter state in *Potash*—is soluble in *Hydrogen Gas*—combines with inflammable substances—and forms alloys with metals.

Potash appears to be a constituent part of most *vegetable* substances; it is also found to enter into the composition of several *minerals*. Usually obtained from the ashes of the former in the state of *common Potash*, by elixation in water and evaporation to dryness; or in that of *pure Potash* by subsequent treatment with *Lime*.

Potash is a part of most vegetable & many
animal bodies. it acts in its pure
state as a caustic. on the addition of
moisture to it heat is evolved. It turns
vegetable blues to green.

exposed to a strong heat it melts & may
be fused into beads. Its power on
substances depends

It is procured from all sorts of timber
& putrefaction. In many woods, Carpenters
have cut down the trees to make potash
the wood is burnt. The Ash of it con-
tain potash. by pouring water on the
gray powder a strong solution of
Potash this is evaporated & which
forms Potash. In the boughs of the
beech & small succulent trees there
is the greatest share of Potash.

many of the weeds in England have Potash
in it common Fennel, Fenitans &c.

were yield Potash by burning. Wormwood
will also yield Potash by the same Process
Pearl Ash is the common Potash purified by
breaking it into small pieces exposed to heat
& the Salt is made red hot... The Oxygen of
the Air takes away the Carbon Hydrogen...
and the substance becomes white forming
the Pearl Ash.

In order to obtain Potash from Tack:
Pearl Ash 1 lb 3 & 4 lbs of Lench. lime
add to them 3 or 4 times their weight of
water Chalk would be formed

we should have a solution of Caustic Potash
this solution being evaporated in a Clean
iron vessel will form the common
Caustic Potash - It is purified by adding
alcohol which has only the property
of destroying the Caustic Potash.

The Oxides of some Metals are soluble in Potash
The Oxide of tin is soluble Oxide of Iron is not.

It has the power of acting on flint &c it will
render them soluble in water...

Struck Potash was thought an elementary substance, not being able to decompose it. But Mr Davy decomposed it by the Voltaic Battery. Potassium is a metal & will amalgamate with other metals alloy it while when put into water decomposes it and renders it alkaline which may be known by dipping a piece of Turmeric paper into it.

100 of Dry Potash contain

Hydrogen Gas may be produced by putting a piece of Potassium into it. It is capable of 3 States of Oxidation. The suboxide which is made by putting a small piece of Potassium & exposing it to the air from which it will absorb oxygen: 2 form Dioxide. 3 Peroxide which is yellow.

Potassium has a very powerful attraction for Oxygen. Put a small bit of Potassium into a retort & heat it by the Air Pump successively and put into the retort some Oxygen which is heated by a lamp of spirit, by this means you will obtain the Peroxide of Potassium. From which comes contact with Potassium will become an alloy & will look like Tin.

Potassium has no action on Ammonia of
Potash.

Potassium with oxygenous Gas.
When heated in Hydrogen it forms an
Hydrosulphide.

Has a strong attraction for *moisture*.

On solution in *Water* produces an increase of temperature (*Liquor Potassæ P. L.*)

Combines with all the *Acids*, and in most instances with a force of attraction superior to that of any other substance.

Fusible in a moderate, and volatile in an intense heat. Is decomposed in a high temperature by *Iron*.

Promotes the fusion of *earthy Bodies* and of *metallic Oxyds*; hence the preparation of different kinds of *Glass*. Unites with *Sulphur*, and renders it soluble in water (*Potassæ Sulphuratum P. L.*)

Dissolves several of the *Metallic Oxyds*. The comparative forces of attraction under all circumstances by no means exactly ascertained.

The order of the attraction of this alkali, in the moist way, *Sulphuric, Nitric, Muriatic, Phosphoric, Fluoric, Oxalic, Tartaric, Arsenic, Succinic, Citric, Lactic, Benzoic, Acetic, Boracic, Sulphureous, Carbonic and Prussic Acid; Water, Unctuous Oils, Sulphur, Metallic Oxyds*; in the dry way, *Phosphoric, Boracic; Arsenic, Sulphuric, Nitric, Muriatic, Fluoric, Succinic, Formic, Lactic, Benzoic, and Acetic Acid, Baryt, Lime, Magnesia, Alumine, Silica, Sulphur*.

Sometimes prescribed internally in solution; in the concrete form frequently employed as a *Caustic*.

Of Soda.

Its *Form, Colour, Taste, Causticity, &c.* nearly the same with those of the former *Alkali*.

Consists, according to Professor Davy, of a peculiar *metallic Base* called *Sodium*, combined with *Oxygen*.

Sodium lighter than water, but heavier than *Potassium*. Other particulars in which these two metals differ.

Found in great abundance in the mineral kingdom, particularly in combination with *Muriatic Acid*.—Obtained, in a pure or caustic form, from *Carbonate of Soda*, by means of *Lime*, as pure or caustic *Potash* is from *Carbonate of Potash*.

Soda has an equally extensive disposition to combine with *Acids* as *Potash*.

Acts powerfully on the *Earths*, and *metallic Oxyds*; also unites readily with *unctuous substances*; hence the preparations of *common Soaps*.

Order of attraction of this *Alkali*, the same as that of *Potash*.

It is impossible to distinguish this from the
fast alkali from its external appearances
it will turn veget. blues green.
It can only be distinguished from Potash by its
chemical combinations. It has a stronger
attraction for acids than ~~Sod~~ Potassium
Oxymuriate of Platinum does not occasion
precipitation in a solution of Soda but
does in a solution of Potap. Soda like
Potap appears to solidify a portion of water
raises the thermometer Soda has been
called fixed or mineral alkali it is diff.
from an animal or veget. substance without a
portion of Soda. It is preserved by burning
some Plants. which is animal it grows in
Mediterranean. It is melted into a sort of
eye Mass which is called Barilla then
Soda Subcarb: in this last substance
it which comes from Spain is best
it is obtained by burning Marine plants
generally there but a small portion of
kali. 100 grains of Acid will saturate 50
grains of Barilla It is used in Manufacturing
it does not contain animal matter

Soda is never found pure; It is found in
Egypt. & on the shore of Persia & other
hot countries

Various Processes have been made to obtain
it from Common Salt. To obtain it
take Sod. Subear. Cl_2 & add Cl_2 of Quick Lime
& Gallons of Water Lime takes the Carb.
Acid. we have a solution
of Caustic Soda which being boiled down
when it is evaporated in the same manner
as you would other Saline Substances
it may be purified by Alcohol -

It decomposes Water & gives off Hydrogen Gas
exactly as Potassium does.

Its Chemical Properties are much the same
as those of the Potassium. - last mentioned
It is capable of 3 States. Suboxide ... 2nd Dioxide
3rd state the peroxide -

25 Oxy 75 Sodium in 100 Parts of Sol
upable of 2 States of Oxidation
Aldehyde & Peroxide

To know the quantity of milk Alkal
in a quantity
weight 100 grams. Lyman
with water & wash
in water & brown. Report
in the alkali. Then take Alkal
in and keep snapping it in
upfully under your hand brought
to the neutral point

2 Ounces of Silver & Peroxide
Bottle. Pour on 2 Gallons of
diluting water, when you are
in a strong solution of Potash
then you separate the water in
the bottle

Ice is quickly Melted by Ammoniacal Gas on account
of its affinity for Water. Ammoniacal Gas is
prepared by a Mixture of ^{2 Parts} Lime & ^{10 Parts} Muriate of Ammonia
and put it over a Spirit Lamp. The dense
vapor which the Muriate Acid gives off
A taper will not burn in Ammoniacal Gas
but sometimes there is a yellow flame.
Ammoniacal Gas may be decomposed by passing
Electric Sparks thro' it. When you wish to
know it very dry Put into it an
inversion of common Dry Bottle

It will combine with all the Acid Gases passing
from the Air form to the Solid State
fill a Phial with Muscated Acid Gas and after
wards you can put a similar quantity of Am-
moniacal Gas without increase of volume
The Ammoniacal Gas has a caustic taste & pungent
& is destructive to Animals made to Respire it
Spirit of Wine Absorbs this Gas as well as Water
The Liquor Ammonia should be stop'd closely -

Of Ammonia.

Form gaseous.—Smell extremely pungent.—
Caustic.—Azotic.—Lighter than Atmospheric
Air—100 cubic inches weigh 18,67 grains—
Experiment of Professor Berselius, shewing its
metallic basis.

Is absorbed both by *Water* and *Alcohol*; by
the former with great rapidity, producing an
increase of temperature and bulk (*Liquor Ammo-
niæ P. L.*) On the contrary, *Ice* dissolved in
this fluid, produces cold.

Is in a slight degree *inflammable*.

Obtained from the distillation of *bones* and
other kinds of *animal Matter*, but purest from
the decomposition of *Muriate of Ammonia*, by
Potash, *Soda*, or *Lime*.

May be decomposed in various ways; as by
exposure to *Heat*, by the *Electric Spark*, in the
reduction of *Metallic Oxyds*, by the distillation
of *Nitrate of Ammonia*, &c. yielding in some
cases *Hydrogen*, in others *Azotic Gas*: which
gasses by particular modes of combination have
been found to reproduce it, and are therefore
considered as its elements.

Agrees nearly with the other *Alkalies* in the
order of its attraction, both in the humid and
dry way.

OF ACIDS.

Appear in general to consist of *Combustible Substances* in union with the base of *Oxygen Gas*, the proportion of which in most cases determines the degree of *Acidity*.—Several of them the immediate result of *Combustion*.

Distinguished by being *sour* to the taste, changing vegetable *blue* colours to *red*, and by their extensive power of combining with other substances.

The properties of the individual *Acids* dependent on the nature of their respective *Bases*: the stronger the attraction of their *Bases* for *Oxygen*, the less intense in most instances their *Acidity*.

All unite readily with water. *Diluted Acids*.

In combination with *Alkalies* form for the most part what are called *Neutral Salts*; with *Earths* and *Metallic Oxyds*, *Earthy* and *Metallic Salts*.

Most of the acids *antiseptic*. Some powerfully *corrosive*.

Admit of different distinctions—from their form, into *solid*, *liquid*, and *gaseous*; from their

The peculiar Smell of Hawks horn arises from
a portion of Empyrenumatic Acid. This makes it diff
from Liqueur Ammonia. Ammonia is contained
in various Vegetables in the ^{Clas} Tetradynamia as
cabbage &c. Ammoniacal Gas may be obtained from
urine. Goat &c To procure it from Animal
Substances a large iron vessel is used by the Man
factories of it when the horns by burning have
become quite white they look as if they contained
nothing but horny substance. The first thing
the Manufacturer does is to break down the horns
and baled, put into the iron vessels from which
the gas is going to them by turn and there again
with Casks which receive the Ammoniacal
Gas

The Filings which has a powerful attraction for Oxygen
makes in it with a little Nitrate of Copper. The Filings
the Iron Filings combine with $\frac{1}{4}$ Nitric Acid.

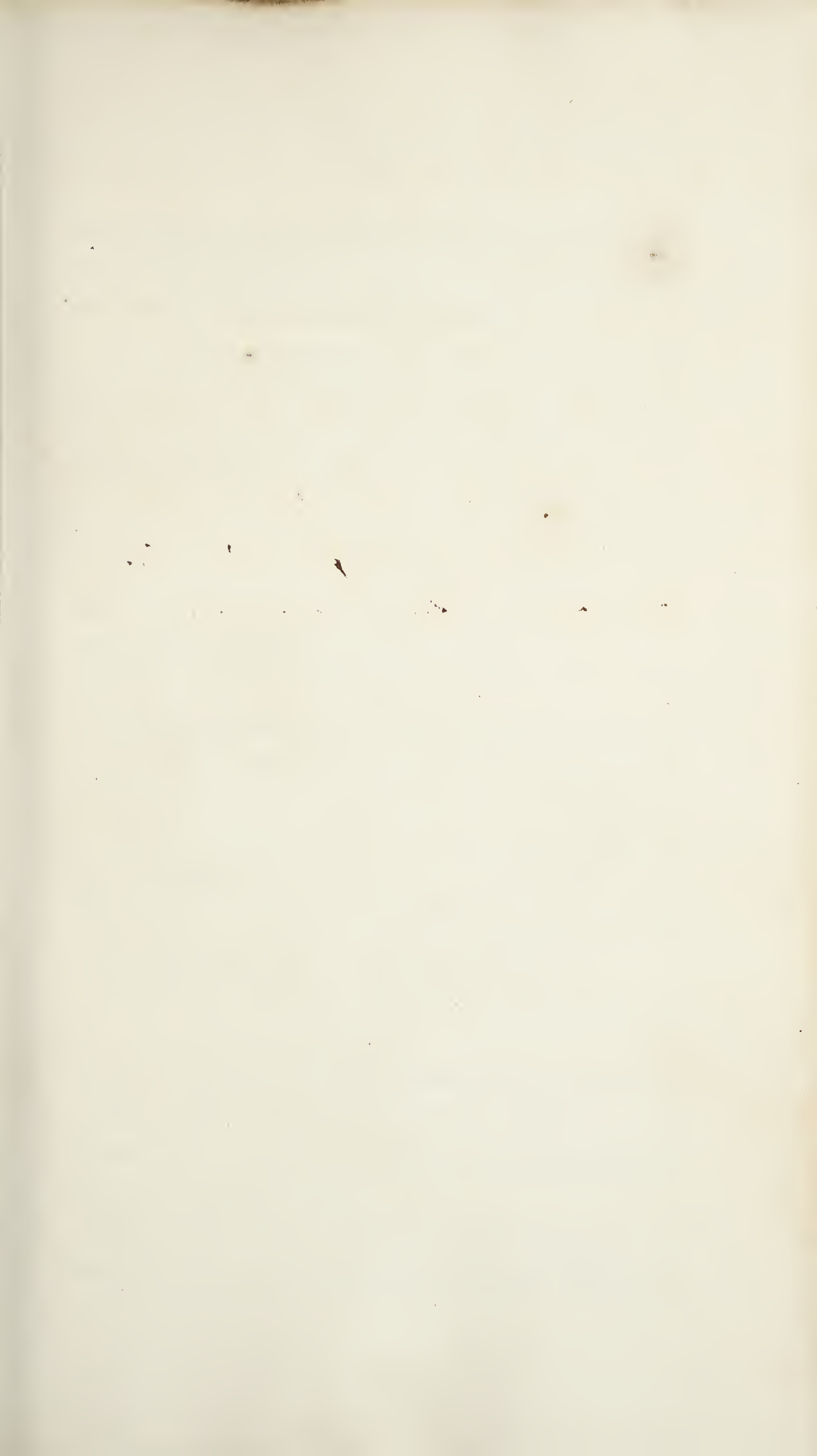
By then using in a little quick lime you decompose

The Best for Ammonia is to maintain a lot of Clafs with
Ammoniac Acid it will produce white fumes.

portion of Ammonia is evolved when urine meets
in the open air. Ammonia combines with
oil & animal & vegetable Principles forming
an imperfect Soap. Eau de lue is made of oil of
Spirits Amm. R^e Meas 4: 3; Oil. Sassa. M^{xxxx} feet grow

Pure Ammonia acts only the Metals when they are in a State of Oxide (that is without you admit Oxygen) Copper put into liq. Ammonia and exposed to the Air it attracts Oxygen and changes its color, but if it is excluded from the Air it is not acted upon

Seberg of Jene in 1800 discovered that Mercury was acted upon in a peculiar manner by subjecting it to Galvanic Influence in a piece of Carb: or Nucleus of Ammonia, an amalgam is formed. The Mercury extending itself & combining with ~~Hydrogen~~ Ammonium



more usual and abundant sources, into *mineral*, *vegetable*, and *animal*; into such as have *simple*, and such as have *compounded* bases.

The more important of the *Acids* are, the *Sulphuric*, *Nitric*, *Muriatic*, *Carbonic*, *Fluoric*, *Boracic*, *Acetous*, *Tartareous*, *Oxalic*, *Gallic*, *Phosphoric* and *Prussic*.

Of Sulphuric Acid.

Formerly called *Acid*, *Oil*, and *Spirit of Vitriol*.

Long considered as the *Universal Acid*.

Seldom found uncombined.

Obtained artificially from the rapid combustion of *Sulphur*, or of a mixture of *Sulphur* with *Nitre*, or from the distillation of *Sulphate of Iron*.

Concentrated afterwards by *boiling*.

When pure, *limpid*, *ponderous*, *unctuous*, *inodorous*, and intensely *sour*.

Specific gravity nearly double that of *Water*.

Has a powerful attraction for *Moisture*, and on mixture with *Water*, occasions a remarkable increase of temperature.

When volatilized by the higher degrees of heat, it assumes the form of a dense white va-

pour, the first portions of which sometimes congeal if exposed to a moderate cold. (*Glacial Oil or Acid of Vitriol.*)

By treatment with *Combustible Substances* is generally more or less discoloured, and may be either deprived of a portion of its *Oxygen*, and thereby made to assume the form of *Gas*, or totally decomposed and reduced to its original basis, *Sulphur*.

The properties of *Sulphureous Acid Gas*, which may be also prepared by the *slow* combustion of *Sulphur*, in many respects different from those of the *common Acid*; its compounds therefore differently denominated. *Sulphite of Potash*, &c.

The *Sulphuric* superior to most *Acids* in its *Power of Attraction* for other bodies.

Independently of its *Water*, supposed to consist of 42,3 parts of *Sulphur* and 57,7 of *Oxygen*.

The order of its attraction *Barytes*, *Strontian*, *Potash*, *Soda*, *Lime*, *Magnesia*, *Ammonia*, *Alumine*, *Metallic Oxyds*, *Water*, *Alcohol*.

Combined with *Alkalies* it forms :

1st, *Sulphate of Potash* (*Potassæ Sulphas P. L.*) usually prepared from the saline mass which remains after the distillation of *Nitric Acid*.—Form, hexangular prisms with hexahedral pyramids—prism frequently wanting.—Taste, *saltish*









bitter.—Difficultly soluble in water; also very difficult of fusion—May be decomposed either in the *moist* way by *Barytes*, or in the *dry* way by calcination with *Charcoal*.—100 parts consist of 42,2 of *Sulphuric Acid*, 50,1 of *Potash*, and 7,7 of *Water* —Application chiefly *medical*.

Super Sulphate of Potash—crystallizes in slender needles or hexangular prisms—soluble in five parts of water, contains twice as much acid as the sulphate. (*Potassæ super Sulphas* P. L.)

2d. *Sulphate of Soda* (*Sodæ Sulphas* P. L.) obtained by solution and crystallization from the matter left behind in the preparation of *Muriatic Acid*, *Muriate of Ammonia*, or *Muriate of Quick-silver*. Is also found native in many mineral springs. Form of its crystals, *hexangular prisms with dihedral summits*.—Taste, *bitter*.—Effloresces on exposure to *Air*.—Readily soluble in *Water*.—When exposed to *Heat*, undergoes watery fusion. May be decomposed in the same way as *Sulphate of Potash*.—100 parts consist of 23,52 *Acid*, 18,48 *Soda*, 58 *Water*.—Used only in *Medicine*.

3d. *Sulphate of Ammonia*, obtained by the union of *Sulphuric Acid* with *Ammonia*, on the addition of diluted *Sulphuric Acid* to liquid *Carbonate of Ammonia*.—Form, *crystalline*.—Taste,

bitter, pungent.—Easily soluble in *Water*. *Fusible*. *Volatile.*—Employed principally in the manufacture of *Muriate of Ammonia*.

Sulphuric Acid forms also with the *Earths* and *Metallic Oxyds*, particular compounds, to be hereafter spoken of under their respective heads.

Employed principally in *Dying*, *Bleaching*, *Tanning*, purification of *Oils*, and in *Medicine*.

Of Nitric Acid.

The basis of this *Acid* is *Azote*—consists of *Nitrous Gas* or *Nitric Oxyd*, and *Oxygen*.

100 parts of *Nitrous Gas* consist of 43 *Azote*, 57 *Oxygen*.—100 cub. inches weigh 33,92 grains. Partly absorbed by *Water*—effects on animals—and on combustible substances—decomposed by the *Electric Spark*—also by *Sulphurets* and some of the *Metals*—means of procuring—test of its purity—reproduces *Nitric Acid* by the addition, and furnishes pure *Azote* by the subtraction of *Oxygen*.—100 parts of dry *Nitric Acid* said to consist of 68,06 *Nitrous Gas* and 31,94 *Oxygen*—or about 29,5 *Azote* and 70,5 *Oxygen*.—*Nitric Acid Gas* rapidly combines with *Water* and

Nitric Acid

In the Nitrous Oxide & Oxygen is in a State of
condensation. It consists of 1 Measure of
Oxygen and half a Measure of Azote

Azote can exist in a diff. State that is it can take more
Oxygen Gas.

1 Measure of Oxygen & Azote form Nitrous Gas

2 Vol.

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Nitrous Gas when in Contact with Oxygen absorbs it
rapidly and forms Nitric Acid. Nitrous Gas is
usually made by Pouring ^{on} Diluted Nitric Acid.

^{Nitrous}
Phosphorus will burn splendidly in Nitric Acid Gas

There has been much diffy of opinion concerning the
quantity of Nitrous Gas & Oxygen to form the
acid. Mr Allen thinks it is about 1/2
of each but a large quantity Oxygen the acid
is diffy.

Nitric acid has diffy to procure as even London
knows in substance to compare. The only way
to get it is to take a glass of water & a pound of
Nitrogen & heat it. It has a very strong
odor for its weight.

It is heavier than Nit. Gas & but it is very
to procure. - It is prepared from 1 Part
of Acid of Sulph. & 2 Parts of Nitric & 2 Parts
The Lute employed on these occasions should
be the fat Lute.

Nitre Acid is made by 60 Parts of Pot. Nitre
29 Parts of Concentrated Sulphuric Acid. Set
Wolfe Apparatus together. In the Process of
Distillation the heat should be very moderate
otherwise when the Salt becomes fluid it will
be thrown out of the retort and perhaps
break the receivers. * you get more Nitric
Acid by a slow Process. — if you wish to
get every portion of Acid you must use a larger
portion of Sulphuric Acid. to the same portion of
Potash Nitre. In large Manufactories they use Iron
Cupels for Preparation of Nitric Acid. This process
will procure it suff: pure for the Dyers & best for Rice
Chemical Experiments it is not fit for it contains
a portion of Acid. Muriat: when by passing water
it — solution of Nitrate of Silver when there
will be a cloud. if there is Sulphuric
Acid: the Nitrate Baryt will discover it by —
cloud being formed. The Acidness in
all these Cases will be Potash Sulphur
if Acid: Nitre: is exposed to the air & light it will
be deprived of part of its Oxygen therefore reduced
to Nitrous Gas. it becomes colored. but it may be
restored by Distillation but this will decompose
it and lessen the quantity. Nitric Acid will
boil when diluted at 219 or 220. but when concentrated
it will not boil till the thermometer rises to

forms *liquid Nitric Acid*.—100 parts spec. grav. 1,5 consist of 91,55 *Acid* and 8,45 *Water*.

Obtained from the decomposition of *Nitrate of Potash* by *Sulphuric Acid*, or by *Sulphate of Iron*.

Purified by re-distilling it from a fresh portion of *Nitre*, or by the addition of *Nitrate of Silver* and of *Baryt*.

In its common form, of a *yellowish* or *orange* colour: when pure, altogether *colourless*.

More *volatile* and less *ponderous* than *Sulphuric Acid*—its *Acidity* equally intense—attracts *Moisture* also from the atmosphere. (*Acidum Nitrosum dilutum P. L.*) *Aqua-fortis*.

Destructive, more immediately than the *Sulphuric Acid*, of the life and texture of bodies to which it is applied.

On mixture with *that Acid* communicates to it the property of dissolving silver: *Aqua Regina*.

Experiments of Cavendish and Milner on *Nitric Acid*—its spontaneous production by processes of nature.

Nitric Acid dissolves all the *Earthy Bodies* except *Silica*.

In its action on the *Metallic Bodies*, it is decomposed to a greater or less degree according

to their capability of uniting with a larger or smaller proportion of its *Oxygen*.

Similar effects produced on it by *Charcoal*, *Phosphorus*, *Sulphur*, and most other combustible substances; as also by exposure either to *Light* or *Heat*.

Hence the difference, if any, between *Nitrous* and *Nitric Acid*, and the formation of *Nitrous Gas*, and *Gaseous Oxyd of Azote*.

In some instances, the reciprocal action such as to occasion immediate inflammation, as on mixture with *Essential Oils*.

Combined with *Alkalies* it forms,

1st. *Nitrate of Potash* (*Potassæ Nitræ P.L.*) Obtained in the East Indies and other countries, by the elixation of certain soils, in which it is either spontaneously and repeatedly produced by their exposure to *Air* and *Light*, or formed upon the addition of *Potash*.—Those soils more especially productive which abound in vegetable and animal matter that has run into a state of putrefaction.—Freed from extraneous saline matter by repeated crystallization.—Form of its crystals *prismatic*.—Taste, *cold, saline, penetrating*.—Much more soluble in hot than in cold water.—In a moderate heat it undergoes watery

any is of opinion when Nitric Acid is combined
with a base the proportion of Oxygen one
and a half.

Nitric Acid is more immediately active on
Ladur than Sulphuric Acid. & when mixed
with the Sulphuric Acid gives the power
of dissolving Silver. Nitric Acid stains
the cuticle when applied to it and destroys
the Camdesh mix^t 5 Part of Oxygen
to 4 parts of Atmosphere Air. When he
introduced a portion of Soap lye
taking there it sparks of electricity
supplying paper with the solution which
sank deep he found it much altered

Nitric formed Nitric Acid from the black
oxide of Manganese & ammoniac. The
powdered part with a Potash solution

Nitric Acid may be decomposed by making a
tube in the middle red hot which has a
strong attraction for Oxygen and this tube
is communicating with a retort in which
there is Nitric Acid. when heat is applied to
the retort Nitrous Acid Gas will be generated.
It has been found if you slowly another portion of Oxygen
is introduced & you will see the Nitrous Oxide gas rise
Nitric Acid may be existing in old rubbish but it may
be decomposed. if you do not know the
quantity of Nitric Acid. The soil on which plants
are sown will give you an idea of the quantity

Charcoal powder when dropped into very
strong Nitric Acid ^{very good} will produce Sparks of
fire. Nitric Acid ^{should} should be never kept in
large quantities for it is a dangerous material.
It is so very combustible: its capacity of taking
up steam & the quantity of it is that it is impossible
to keep it in vessels or Nitric Acid from being
exposed to heat then from heat.
Essential Oils will be immediately decomposed
by Nitric Acid: as Oil of Turpentine: the other
Oils consist of Carbon & Hydrogen & are
transformed by Nitric Acid into
Nitric Acid will decompose all the earths except
Silica

you may always know a Nitrate by heating
it it will yield Oxygen

Nitrate of Potash when ignited as a gas
but of course Carbon it is a very common
a substance

The quantity of Gas produced is very dependent
on the Character of the substance which the Acid
will be decomposed.

In Purifying Nitrate of Potash you must take
care not to apply the heat too long but just be-
fore it runs down in water & forms a clear
solution.

Melted Nitrate of Potash is an excellent test
for Carbonaceous Matter. Take a little of it
and pour it on Charcoal when it will exhibit
a pleasing Phenomenon, arising from its
strong attraction for Oxygen. By a combi-
nation of these two Substances Gunpowder
is made. The parts 75 Salt Petre, 15 Charcoal, Sulphur
Parts. Salt Petre is fused. Charcoal should be very good
is best made in Iron Cylinders not in the Common Method.
Nitrate of Potash may be decomposed by Sulphur
& Metals as well as by Charcoal.

Being provided pure Ingredients. The Salt Petre
Charcoal & Sulphur: is all ground to a fine
powder (but separately) they are all mix'd
in a Vertical Stam working on an horizontal
one - If by accident it explodes at the
beginning of the Process it will not explode
very violently, but soon so when the pro-
cess is nearly finished. When thus
prepared it is pressed into a cake by
a press, it is then broke into small
pieces - and sifted into pieces of the size
of Common grains of Gunpowder. These grains
are dried afterwards: it is some-
times dried by Steam. It is before

When Gunpowder is good it will explode
from off paper without making any mark
on the paper. -

12 Grains of Gunpowder in exploding will lift
a very considerable weight 2000 Pounds
Gunpowder exploded in a proper apparatus
for demonstration evolution of Carbon Dioxide & Azote
very little water formed on the explosion
of Gunpowder

Nitric Acid: Combined with Soda forms
Nitrate of Soda & it resembles very much
Nitrate of Potash if Crystallized in Carbon

Nitric Acid Combined with Ammonia forms
Nitrate of Ammonia:

when Crystallized it is a six sided Prism

Soluble in twice its weight of Cold water
& half its weight in boiling water

Nitrate of Ammon: should never be kept in
Metallic vessels: 1 part of the Salt and
2 parts of Nitric Acid: will form a fluid
Capable of dissolving Gold.

fusion (*Salprunellæ*).—An increase of the heat produces a decomposition of its *Acid*, the *Alkali* remaining unchanged.—May be more rapidly decomposed by the addition of *Charcoal*; hence the preparation of *Gun-powder*.—When decomposed in the moist way, by treatment with the *Sulphuric*, furnishes *Nitric Acid*.—100 parts dried at 70 *Farenh.* consist of 44 of *Nitric Acid*, 51.8 of *Potash*, and 4.2 of *Water*.—Used extensively in *Glass-making*, *Metallurgy*, preparation of *Gun-powder*, and for *dietetic* and *medical* purposes.

2d. *Nitrate of Soda*, called formerly *cubical* or *quadrangular Nitre*. In most of its properties resembles the former.

3d. *Nitrate of Ammonia*; obtained by slow evaporation, from the combination of *Nitric Acid* and *Ammonia*.—Form, *crystalline*.—Taste, *cool*, *bitter*, *urinous*.—*Deliquescent*.—Easily *fusible*.—Under cautious distillation, yields *Gaseous Oxyd of Azote*, *Nitrous Gas*, and *Water*; but detonates when suddenly heated.

The *Order of Attraction* of this acid, the same as that of the *Sulphuric*.

Chief uses, in *Dying*, *Etching*, and *Assaying*; and sometimes employed in *Medicine*.

Of Muriatic Acid.

Obtained by decomposing *Muriate of Soda*, by means similar to those employed in the preparation of the *Nitric Acid*.

Its purest form *Gaseous*.

In this state heavier than *Atmospherical Air*; 100 cubic inches at mean pressure and temperature, weighing 39 grains; of a pungent *odour*; *irrespirable*; destructive to *Flame*, imparting to it under extinction a bright green colour; inflames the skin without discolouring or corroding it.

Formerly supposed to consist of an unknown radical intimately combined with *Oxygen*. Late researches of Professor Davy, by which this doctrine is controverted.

When dry has no action on the *Metals*, nor on any other of the *Combustible Substances*.

Exposed to a moist atmosphere it becomes *cloudy*—Is readily absorbed both by *Water* and *Ice*, the latter of which it liquefies: in either case it occasions an increase of *Temperature* and augmentation of *Bulk*, communicating to the water the general properties of an *Acid* (*Acidum Muriaticum* P. L.)

Muriatic Acid

is one of the component parts of Muriatic Acid -

many attempts have been made to decompose it. It is obtained from Salt by Sulphuric Acid, in order to preserve caught up Pounds of Common Salt put it into water & connected with Woolfs Apparatus: then Put in Pounds of Water Pounds of Concentrated Sulphuric Acid: which is put into the Retort with the Muriate of Soda...

This way you may procure tolerably pure. The Muriatic Acid in large Manufacturers is made in large iron vessels & is not so very pure for their purposes but it is not good enough for Chemical purposes. Wh. Muriatic Acid contains Iron it is usually a brown color after it takes up: as much as 400 times bulk of Muriatic Acid. Gas

The Muriatic Acid in its purest form is in $\frac{1}{4}$ state of Gas.

This gas extinguishes a taper & unfit for respiration You procure this Gas by taking 2 parts of Cam: Salt heat it you then put this Salt into a retort & then pouring on it a little Sulphuric Acid: and apply a gentle heat when you may by means of a bladder collect this Gas. —

This Gas when applied to combustible Substances does not have a very considerable effect.

The liquid Muriatic Acid will readily dissolve most of the Metals especially if it is exposed to the Air, it readily melts Zinc and Iron but it has not any action on Gold Platinum. The liquid Muriatic Acid combines readily with $\frac{1}{4}$ Alkalies and most of the Earths.

A solution of Nitrate of Silver is a good test for this Acid, by making a white precipitate.

When Muriatic Acid is mixed with Nitrate of Potash it forms the Salt of Silver.

matic and contract on Gold Platinum Molybdenum

minerals are decomposed by Nitric & Sulph: A

withy Minerals are particularly deliquescent
so much moisture from $\frac{1}{4}$ air.

argent: & it: is a very delicate test of the presence
of silver: it decomposes it by causing a
white Cloud. & it is also enables us to
exactly how much silver there is. Nitrate
is formed on the one hand & Minera-
lized on the other or Lime Carbon by the old
which contains 19 Parts of Acid
oxide of Silver

its weight
mineral Salt soluble in 3 times of Water
very little more if heated. Alcohol dissolves
our common Salts. if Pure (Spirits) it
is less than 1/2 grain 0.15 oil 0.20 Water
1000 The Minerals of Soda are very
rare. etc. Crystals are cubical various
crystals don't deliquesce when Pure
are, but it is usually containing water
which causes it to deliquesce generally
in Sulph: of Magnesia & Lime this causes
Part of Sea Water then salt form
incredible portions Diff: proportions
of parts At the Baltic is estimated Salt

Common Salt generally has in it
Sulphate of Magnesia & Sulphate of Lime

It constitutes the Salts of the Sea Water
it may be procured by evaporation
Muriate of Soda found in immense quantities
in differ parts of the World as Cordova &c
Rafra in some places so abundant & low
that they build huts with it. In the
Christian it is found. generally covered
by a mixture of argill &c
for preparing it a certain place is chosen
by the side of the Sea or Pond is made
and by the side of this pond there is another
by and there as third near inland
these ponds are exposed to the heat of
the Sun which of course is covered off
partly by evaporation: it is then let
into the 2nd Pond: & then into the third
which is nearest the boiling house
when it is let in and boiled undevapor-
ated & Chrysallization soon commences. it is
then put into baskets thus called Basket Salt
and another form of it is the Bay Salt

Muriatic Acid dissolves also some of the *Metals*, as *Zinc* and *Iron*, with the production of *Hydrogen Gas*; and shews a superiority of attraction for most of the *Metallic Oxyds*.

In its liquid and more common form it readily dissolves and unites with all the *Alkalies* and most of the *Earths*.—Its combinations with the former are,

1st. *Muriate of Potash*, formerly called *Salt of Silvius*; a salt but little used.

2d. *Muriate of Soda (Common Salt)*; obtained by evaporation from *Sea Water*, or the water of *salt Springs*, or the solution of *Rock Salt*, which is found in several parts of the world in immense quantities.—Form of its crystals *cubical*.—Taste, agreeably *saline*.—Equally soluble in cold *Water* as in hot; soluble also in *Alcohol*.—Crystals burst or decrepitate on sudden exposure to *Heat*. Melt in a red heat without decomposition. May be converted into vapour by being intensely heated.—Promote the fusion of many of the *earthy* and *metallic* bodies.—May be decomposed in the moist way by *Sulphuric* and *Nitric Acids*; and in the dry way, by the *Phosphoric*, *Boracic*, and *Arsenic*. May also be decomposed by *Oxyd of Lead*; hence the preparation of *Turner's Patent Yellow*.—100 parts *Mur.* of *Soda* dried

at a red heat, consist of 46 *Mur. Acid*, and 54 *Soda*. New views of Professor Davy on the constitution of this *Salt*—Other methods of procuring *Soda* from it—Of extensive application in *Agriculture*, *Glass-making*, *Glazing*, *Metal-lurgy*, *Soap-making*, *Diet*, *Pharmaceutic Chemistry*, &c.

3d. *Muriate of Ammonia*, (*Ammoniacæ Murias. P. L.*) ; found native in the neighbourhood of *Volcanos* ; prepared also, in large quantities, in the dry way, by double chemical affinity from a mixture of *Sulphate of Ammonia* and *Muriate of Soda*.—Form, *concrete*.—Taste, *penetrating, acrid, urinous*.—Soluble both in *Water* and *Alcohol*.—Crystallizes under evaporation into small quadrangular *Prisms*.—Yields its basis readily to both the fixed alkalies, and to lime :—is also partially decomposed by sublimation with oxyd of iron. (*Ferrum Ammoniatum P. L.*)—Of extensive application in *Dying*, *Tinning*, *Soldering*, &c. Used also frequently in *Medicine*.

The chemical affinity of the common form of the *Muriatic Acid* in most instances weaker than that of either the *Sulphuric* or *Nitric* ; but the order of its attraction the same.

On mixture, either in the liquid or gaseous state, with *Nitric Acid*, occasions in it a partial

Scotch Salt is not so Pure

The Sulphate of Lime in Sea Water embarrasses
the makers of Murate of Soda extremely

The Method of Decomposing Common Salt is
effected by Diff: Methods.

One has 7 property of Decomposing Common
Salt by mixing it into a paste and putting
to a damp Cellar - ^{in an iron} Discharge which is an
oxide of lead will also decompose ^{4 Pounds of 1-Pound of C. Salt} forming
Color Called Turner's yellow - one may likewise
decompose it by Barytes but one must be
collect that this the Barytes is poisonous
urate of Ammonia is found in Volcanos
The Mountains...

so in Animal fluids usually met with.
a Concrete for or taking the ~~use~~ shreds of 4
lb. The Taste Urinous, Acid: Much used
Soldering, and Dyeing. in Soldering two pieces
Metal it is necessary that both surfaces
should be very bright that is free
from Oxide

There is a salt permeation from the Bay
salt which is formed by long exposure to
the air & Sun & Chry, falling slowly on
the Ponds. It is thought to contain more
superiority than the other. If its better
actual superiority depends on the largeness
of the grains which decompose by rapidly
thus forming the most perfect completion.

What they call in Cheshire the Sham Salt
is made by evaporating the water and
hastily the resulting Salt is called by the
Sham Salt.

For the purposes of the large grain
salt they only raise the permeation to
degrees & let it continue a long time
During the evaporation of the Salt the
crystals form a pile for and which
will be found cubical. &
gather together & ultimately form to
bottom

Exp of Am allum
Anhydrous Salt is made by depositing
Salt in Sea Water & forming
is contained 99th of Pure Muriate of Soda in
Wilmington Salt contained in 100
parts ~~contains~~ 96 parts of Muriate
of Soda

in the Mother Ligner they make
Whack of Magnesia

Soda is so very useful it has been
objects to obtain it from Common

Duty on Common ^{Salt} is enormous. but
inventions take off this Duty when
it is used to make Soda.

Muriate of Soda is useful in fluxing
Metals.

Muriate of Potash but little used.
more soluble in hot than cold water
Muriate of Ammonia consists of Acid
of Ammonia: usually met with
the in concrete form very
soluble in water & if carefully
purified may be made to crys-
tallize in very pure form

The Oxymuriatic Gas was first formed by Shal
when you present a Subst: containing Oxygen to Heat &
Supply heat you obtain a peculiar Gas. Chlorine
This considered as a simple substance, 1774. Davy made the disc-
Take 3 parts of liquid Muriatic Acid, and put
into a retort, with black Oxide of Manganese 10
Water capable of dissolving twice its volume of the
Gas - then form what has been called liquid ^{Ox} Muriatic
acid

~~Liquid~~ Oxymuriatic Acid will become decompo-
sed by light, but not by air or heat.
The liquid Oxymuriatic Acid is sometimes given in Medicine
it is very much used in Bleaching. ~~but~~ Speedily de-
the Gum color of vegetable in fact all colors except
yellows.

Muriatic Acid will weaken vegetable fibres
exceedingly if the solution is too strong. The Manner
make use of Tim with it, then called Oxymuriatic of Tim
The smell of Oxymuriated Muriatic Gas is pungent
not respirable, 100 Cubic Inches of it weighs 70 Grs

it may be procured from oxymuriate of Potash
and strong Muriatic Acid. a very gentle
heat must be used in obtaining this Gas by these
means

This Gas has an extreme effect on vegetable Colors
it acts powerfully on Putrid Infection, but from
the suffocating Nature of it, you must not suffer much
to be evolved: Take care to remove every portion
of Gas from the room for it will spoil it
Muriate of Soda Manganese & Diluted
Sulphuric Acid in a Lignum will give it
and it can be brought out in a dark room
in a dark room

considered Chlorine which he named it
the new acid gas
100 Cubic Inches weight 75 Grs
burns in it with a very luminous flame
adopt Davy's theory in 1808 suppose that

33

his was Chlorine

decomposition, and acquires new properties:

Aqua Regia.

An analogous change found to be produced
on it by distillation from certain of the Metallic
Oxyds, more especially from that of Manganese:
hence the preparation of Oxy-muriatic, or as it
was at first called, Dephlogisticated Marine
Acid.

New views of Professor Davy on the nature
of this Gas—Experiments tending to prove that
it is a simple substance—and if so, the present
name improper.

The sensible properties of this Gas distin-
guishable from those of all the other Elastic
Fluids—Remarkable for its effects both on Ani-
mal and Vegetable Colours—Thought to possess
the important property of destroying putrid and
contagious Effluvia—Not so readily absorbed by
Water as common Muriatic Acid Gas—Is mate-
rially affected by exposure to Light, is readily
acted upon by most of the Metallic Bodies, and,
in several instances, produces appearances like
actual Inflammation.

Much heavier than Atmospheric Air.—100
cub. inches weighing 74,5 grs.

Fin Put into Chlorine makes divaric
any degree. Calomel: & Oxymuriatic of
any gas. Compound of Muriatic
Chlorine. If a little Sulphur of Antimony
put into Chlorine and a little Spark
it will give off a flame which is

When equal parts of Chlorine & Hydrogen are mixed & then Hydrogen is present it is formed as before
the chlorine gas is added to it in the same way it is formed as before

take fire in the Gas without having been
commonly heated Davy says Phosphor-
ous combines with Chlorine in 2 diff^t p^{ts}
forming 34 2 diff^t Substances the
Phosphoric & Phosphorous the last is a fluid

Ideas of Scheele on the constitution of this
singular substance—Of Berthollet and the
French Chemists—Of Dr. Henry—Important
discoveries of Professor Davy; reasons which
induced him to doubt the existence of *Oxygen* in
it—Curious effects on *Phosphorus*—Highly ne-
gative with respect to *Electrical energy*—Means
of procuring.—Singular compound of this Gas
and *Oxygen*.

Its combinations with salifiable bases (hitherto
called *Oxy-muriates*) afford *Oxygen Gas* when
heated; triturated with combustible substances,
they inflame and frequently detonate. The most
important of these is, *Oxy-muriate of Potash*,
obtained by cautiously evaporating and cool-
ing a solution of *Potash*, saturated with *Oxy-*
muriatic Acid. It crystallizes into flat hexhedral
prisms or plates, of a pearly *lustre*: *Taste* faintly
saline: moderately soluble in *Water*: on expo-
sure to *heat* readily undergoes watery fusion,
and, like *Nitrate of Potash*, yields great abun-
dance of *Oxygen Gas*, but of more than ordinary
purity.—Much more remarkable than *Nitrate of*
Potash, for its power of detonating on mixture
with several of the combustible bodies, as *Char-*
coal, *Sulphur*, and *Phosphorus*; more especially
the latter.

Mixture of Potash & Sulphur. And is good & for
Kilnmen uses it as an infermentary Conlage

Hydrochloric Acid Gas acts powerfully on
Surface of Mercury

If you take pure Hydrochloric Acid Gas, heat
Potassium in it, it will take fire after
being well heated, you put the Potassium
to a retort & exhaust it, by means of the
Air Pump you then fill it with Hydrochloric
Acid Gas, when by applying a lamp to the
tub you will find the Potassium burn
vigorously. The Action of ^{Oxy} Hydrochloric Acid Gas
Chlorine and the Potassium is very obvious
has highly negative properties. When next
the Hydrogen Gas a remarkable change takes
place. Take equal portions of Hydrogen &
Chlorine it will be found when there are
it common Hydrochloric Acid, this change
directly brought about by exposing a mixture
of these 2 gases to the light of a Sun it will
ignite. Phosphorus will combine with
Chlorine or Hydrogen. when you introduce
Chlorine or Phosphorus it will take fire
if done in a retort.

It will act very Powerfully on Sulphur also.
When Mixt with Carbonic Acid Gas forms a peculiar
Compound, it forms a gas heavier than any
other Gas.

In this Compound the Oxygen is certainly
Condensed

The Smell of this is different to Chlorine
it does not act so powerfully on Metals
as the Chlorine.

The Bleaching properties of Chlorine, is obvious
it will change a piece of Paisley almost
white. (Chlorine is also called Oxymuriatic Acid
Gas)

Flames of Sulphur exposed to Chlorine
the Tin is melted in it about 7 in the
l. of Gas even disappears & what is
is hydrogen

a small quantity of Hyperoxygenate
Pot. & dilute Muriatic Gas & apply
gentle heat you get ~~the~~ Chlorine
immediately absorbed by water
in Chlorine & turns into Bells
of water This Gas is easily separated
from Chlorine by running
the Chlorine gas through
the ~~the~~ Chlorine
This Gas Delanates when
it is applied
Hyperoxy. of Potash is made as very

ingiven 16 or 18 times its weight of water
when it is

alt produces violent effects on being rubbed
Sulphur If this salt is mixt up with Sugar Honey
or any let them on fire by Sulphuric Acid the
retainers from boxes are made in this way

Experiments on the constitution of this *Acid*.
—100 parts by weight consist of 28,60 *Carbon*,
71,40 *Oxygen*.

Charcoal, not an *Oxyd of Diamond* as erroneously supposed.—*Carbon* can exist in two distinct states of *Oxydation*.—*Gaseous Oxyd of Carbon*.

Procured by exposing the *Oxyds of Metals* and *Charcoal*, or a mixture of powdered *Marble* and filings of *Zinc* to a red heat.

Formerly considered as *Hydrogen Gas* holding *Carbon* in solution, and called by *Priestly Heavy Inflammable Air*, but proved by *Cruikshank* to be a distinct species, having *Carbon* for its basis combined with a limited proportion of *Oxygen*, inferior to that which constitutes *Carbonic Acid*.—100 cubical inches, middle temperature and pressure, weigh about 20 grs.—It is inflammable, and burns with a lambent blue flame, but does not explode with common air. And by combustion is converted into *Carbonic Acid*.

This acid, like others, is disposed to combine with salifiable bases. With *Alkalies* it forms,

1st. *Carbonate of Potash* (*Potassæ sub-carbonas* P. L.) usually obtained from the ashes of vegetables, by lixiviating them in water, decanting

the Water Gas is taken in solution of
bicarbonate of Soda

Water Gas is made by passing
the Sulphur or Petroleum or Chalk down
the water employing a burner which
the Gas sent into the water it is driven up
in syringes & by a Piston it is forced into
a very Tube & by this means the Water is
being evaporated

In Lempire (Water) 183 it was observed
Carbonic Acid Gas is taken up
in than water.

When in a proportion with Potash
may the Subcarbonate of Potash

is met with carbonate of Potash in
the States. ²¹ as a Subcarbonate or Salt
Gastar. Pearl ash is like the Subcarbonate
Potash, it may be easily prepared from
Carbonate of Potash.

Carbonic Acid resembles the pector of some acid
with 1/2 of Sulphuric Acid
with 1/2 of Atmospheric air or any other gas

Let Powder Zinc, Marble Powder & Soda
Lime mixed put in a glass vessel &
apply a moderate heat, when the Pure
Gaseous Oxide of Carbon will come

If 2 Vol of this Gas (Gaseous Oxide of Carb) &
volume of Oxygen be mixed & the Electric
Spark there is you will diminish it to 2 parts. which
parts will be Carbonic Acid known by rendering it Opaque
100 Parts the Gas (Gas. Oxide of Carb) consists
of 3 Carbon & 5 1/2 Oxygen

at a Temperature high perhaps Carbon Acid
may be decomposed by strong Acids
forming a strong solution for Oxygen

Some Water is a good Test for Carbonic
acid Gas

Spring Water contains Carbonic Acid

Carbonate of Soda is found at Beni-Suef
and on the coast of Egypt and elsewhere.
It generally prevails in the country
bordering the Ashes of Boullon. It is
very soluble. Effluvescences are exposed
in day atmosphere more particularly

Potash is saturated with Carbonic Acid
and gives a much larger dose of it.

Thollet. Take Subcarbonate of Potash & Soda
and of Carbonate of Ammonia the
of Carbonic Acid will be given over
the Carbon of Ammonia.

Carbonate of Soda may be changed into Carbonic Acid
by quantity of Carbonic Acid

Take Sal Am. 1 Pound 2 Parts
Common Chalk

Carbonate of Ammonia is prepared
from Ammonia. Soluble in 2 Weight
of water. 100^{lb} Consist of 50 Carb. Ac.
10 Ammon. 20 Water. Substantially of ammonia
It is the basis of the Ammonia

In order to fill Smelling Bottles Take Part of
Ammonia Chalk & Part put into a vessel & apply heat. The
heat you see the Ammonia & form a mass of Ammonia & the Carb.
acid & Ammonia, will be combined.

It is Soluble in Water in hence the
preparation of it. Ammonia it
is not Soluble in Spirit

Carbonic Acid is a powerful
Antiseptic. It is employed to foul
Wells as it is when Charcoal is applied

and evaporating to dryness.—Form, *powdery*.—Taste, *urinous*.—Colour, *pearly-white*.—*Deliquescent* in a moist atmosphere (*Liquor Potassæ sub-carbonatis* P. L.) May then be crystallized after cautious evaporation—May be decomposed in the moist way by most of the other *Acids*, or by *Lime* ; and in the dry way, by the simple application of *Heat*.—Uses various, as in *Glass-making*, *Bleaching*, *Metallurgy*, *Medicine*, &c.

2d. *Carbonate of Soda*, (*Sodæ sub-carbonas* P. L.) found native in *Egypt*, the island of *Teneriffe*, and elsewhere ; obtained also from the ashes of certain marine plants, by elixation, evaporation, and crystallization.—Form of its crystals, *octahedral*.—Effloresces on exposure to *Air*.—Its other properties and uses nearly similar to those of *Carbonate of Potash*.

3d. *Carbonate of Ammonia* (*Ammoniacæ Carbonas* P. L.) ; obtained by distillation from most animal, and some vegetable and mineral substances : or from the decomposition of *Muriate of Ammonia* by *Carbonate of Potash*, or by *Lime*.—Form, *concrete*.—Smell, *pungent*.—Taste, *urinous*.—Very soluble in *Water* (*Liquor Ammoniacæ Carbonatis* P. L.) Soluble also in *Spirit* (*Spiritus Ammoniacæ* P. L.) With unctuous substances forms an imperfect *Soap* (*Linimentum Ammo,*

niæ P. L.). Like all the foregoing compounds, may be decomposed by *Potash*, *Soda*, *Barytes*, or *Lime*.

Carbonic Acid now universally considered as consisting of *Carbon* rendered acid by *Oxygen*; for which it has a stronger affinity than any other acidifiable base.

Inferior to most of the *Acids* in its *Attraction* for other bodies.

Order of attraction, *Barytes*, *Strontian*, *Lime*, *Potash*, *Soda*, *Magnesia*, *Ammonia*, *Alumine*, *Metallic Oxyds*, *Water*, *Alcohol*. Uses chiefly *medical*.

Of Fluoric Acid.

Obtained by decomposing *Fluate* of *Lime* by means of the *Sulphuric Acid*.

Form *gaseous*.

Heavier than atmospherical air, *Caustic*.—
Kills animals immersed in it.

Unites readily with *Water*, and renders it intensely sour.

In union with the *Alkalies*, forms compounds of a gelatinous consistence.

Possesses the remarkable property of dissolving and volatilizing *Silica*.

is with dense white fumes, it is also
D in many cases. It is

Fluoric Acid

is by itself a distinct substance, and is not known to be

The Radical for this Acid is not known

is obtained by taking 1 Part of Fluato of Lime

and 2 parts of Concentrated Sulphuric Acid

When we want it in a very concen-

trated form we use a different Method
in part no water in the second vessel

This Acid Corrodes Animal Substances very
quickly therefore we should be very careful
to prevent the Gas arising from the Concentrated
Fluoric Acid.

When mixed with Soda it forms Fluato of
Soda which neither deliquesces or effloresces
when exposed to the air.

It is very readily

This Acid does not act on Lead Silver &c
but when in the state of Oxide it will
act on them very sensibly. ^{It forms Fluorides.} It readily

This Acid effects Glass very sensibly

Coat $\frac{1}{2}$ Glass with Wax and then with a engraving
tool engrave what you want? ... useful in ...
... be hardening and even when also used for

It is used very much in etching of Glass

Water absorbs a large quantity of the Silicate
Fluoric Acid Gas free 200 times its bulk
You can get the Silicate Fluoric Acid Gas by mixing ...
... with the Acid. & fluoride of Lime

On burning Potash in Fluoric Acid Gas
a great portion of Hydrogen is formed.

The Fluoride of Lead is decomposed

Fluoric Acid is capable of combining with Potash
and is combined as much Potash

Fluoric Acid has a disposition to form a
compound with Silver & Platinum Acid.

1 Part of Nitrous Peroxide
Parts of Fluoric Acid, 1 part of Potash
12 times the weight of Acid Sulphur
acid - measures of gas, of which 1000 contains
the weight ⁷³ of hydrogen ~~gas~~ this is the gas which is
used to ascertain the quantity of hydrogen
~~the above is not correct as the gas is not pure~~
the oxygen

Fluoric Acid has not been affected by passing
over red hot iron

Boracic Acid

Comes to us from the East

This was formerly called the *Acidum*
boracicum supposed to be of that kind
Salt. it presents itself in a solid

form and it feels like *tetrum*

Davy found the *Burning* *vitrified* *potash* & Boracic Acid
that he got a Chocolate colored mass

and which being *heated* in Oxygen gas
he obtained again Boracic Acid it will take
up a great deal of water & become a

You procure Boracic Acid by taking *in water & salts*
a solution of Borax, and dropping into

it a little Sulphuric Acid or Phosphoric
until you have saturated it completely
then it crystallizes in white crystals of Boracic

The Boracic Acid requires a great portion
of water of solution it is soluble in water

the solution is colorless & has a
slight alkaline taste & a yellowish

color when exposed to the air it is changed by
the heat of the sun & the heat of the
fire & is very of the acid

Dissolves *Zinc, Iron, and Copper*; the other metals not acted on, unless in the state of *Oxyds*.

Decomposed by Professor Davy by means of *Potassium*.

Order of attraction, *Lime, Barytes, Strontian, Magnesia, Potash, Soda, Ammonia, Alumine, Metallic Oxyds, Water, Alcohol*.

Use—*Etching on Glass*.

Of Boracic Acid.

Called formerly *Sedative Salt*.

Form, *concrete, scaly*.—*Semi-transparent* and of a *pearly lustre*, and *saline acid taste*.

Sparingly soluble in Water; more so in *Alcohol*, to the flame of which it communicates a *greenish tinge*.

When united with *Water* easily sublimed.

Fusible without addition into a transparent *Glass*.

Decomposed by *voltaic Electricity*, but more completely by *Potassium*—Experiments of Davy on this subject—Conjectures that the basis is *metallic*.

By combination with *Soda*, produces *Borate of Soda* (*Sodæ Boras. P. L.*) ; obtained, in an impure form, from the spontaneous evaporation of the water of certain lakes in the kingdom of *Thibet* ; said also to be found in those of *South America*.—Purified by subsequent solution, boiling, and crystallization.—Form of its crystals, *prismatic*.—100 parts consist of about 34 *Acid*, 19 *Soda*, 47 *Water*.—Taste, *styptic*.—Readily soluble in *hot Water*.—Changes vegetable *blue* colours to *green*.—When heated, parts with its water of crystallization, and melts into a transparent *Glass*, which effloresces on exposure, and may be again dissolved in water.—Serves as a flux for all the *Earths* and most of the *Metallic Oxyds*.—May be decomposed in the moist way, by the *Sulphuric*, and several other acids ; in the dry way, by the *Phosphoric* only.—Employed chiefly in *Metallurgy*, and *Glass-making*.

This *Acid* is found in solution in the water of certain lakes in *Tuscany* ; or obtained artificially by decomposing *Borate of Soda* by *Sulphuric*, *Nitric*, or *Muriatic Acid* ; or by subliming it with *Sulphate of Iron*.

Order of attraction, *Lime*, *Barytes*, *Strontian*, *Magnesia*, *Potash*, *Soda*, *Ammonia*, *Alumine*, *Metallic Oxyds*, *Water*, *Alcohol*.

Solution of Boracic Acid in Spirit
When given a green flame when
put on Paper. This flame is
assisted by the blue flame which the
spirit gives, and the other colored flame
which the Borax gives.

When Boracic acid is strongly heated
it becomes vitrified.

It is a cheap base when the Boracic
acid contains a large quantity of the
same acid.

To obtain Borax - heat the Spirit to redness
and pour on Salts. It is separated and
the green spirit obtained. Long. 1800
1800.

The Method of Purifying Borax

Borax is capable of decomposing most of the acids as it has been decomposed them as

Borax does not require an account of its action
in the manufacture of glass

Borax being heated to drive off the
water of crystallization forms glass of Borax

Borax acid is found everywhere abundant
in the form of minerals

The Tartaric acid is found everywhere 30 p. 100
of the acid

Acetic Acid

Vegetable acids have generally composed Radicals
again is Carbon & Hydrogen

kind of them by den them by fermentation
now may be also converted into Acetic Acid
again

In process of changing them appears to be a dispo-
sitionment of particles & an addition of Oxygen
can appear the degree to the acid bodies
being stronger. When vinegar has been
some time decomposed Potatoes
due to keep vinegar is to put it into glass
jar & Boil them in Water & Cork it up
again as before to contain great deal of Vinegar
Acid as Oxalic Acid Tartaric Acid
this Humidity which gives common vinegar
its color. For internal purposes carry on the
action in Glass vessels

Specific Gravity of distilled vinegar is
that of Water

very small quantities of Potash may be
readily concentrated

this process can be conducted in by evaporating
action of the boiling water. Potash. If you have
much of Potash with much Acetic Acid you will have
a great deal of Potash which can be found

in the same manner Acetic Acid combines with
Hydrogen of Metals in Silver Copper with the com-
mon acids. Distilled Vinegar is nothing more
than common Vinegar boiled down & evaporated
you obtain a beautiful clear liquid.

Vinegar may be prevented from undergoing
a change to Putrefaction, by having $\frac{2}{3}$
Vinegar in Bottles, and place these Bottles
in boiling Water and then Cook it, and
it will keep for a long time.

For the Purification of Vinegar for internal
purposes, we must not use Metallic Vessels.
We must use Glass for this purpose.

Common Vinegar is very much diluted
so that we can take it in our food by
a common Spoon.

We can procure strong Acetic Acid by
adding to the Vinegar Potash an effervescence
takes place, when $\frac{1}{2}$ Particles of
Potash combine with $\frac{1}{2}$ particles of strong
Acetic Acid, the result of this process is $\frac{1}{2}$
Acetate of Potash, by evaporation you obtain
this salt if too great a heat is employed
it will be broken and will not be soluble
in Alcohol or Water entirely.

Of Acetic Acid.

Procured from a number of substances capable of undergoing the vinous fermentation—Is largely diluted with *Water* in common *Vinegar*, and may be obtained from it by cautious distillation—Procured also by distilling *Wood*—When pure, without colour, and in taste moderately acid—May be concentrated by freezing—But most perfectly by combining it with a *salifiable base* and subjecting the compound to distillation—As *Verdigris*, or *Acetate of Potash* and *Sulphuric Acid*—Other methods—Sometimes called *Radical Vinegar*—Very *acrid*—Inflames the skin—Slightly inflammable—Thought to consist of 50 *Oxygen*, 14 *Hydrogen*, 36 *Carbon*—*Aromatic Essence of Vinegar* combined with *Potash* produces *Acetate of Potash* (*Potassæ Acetas P. L.*); prepared by saturating *Potash* with *distilled Vinegar*, evaporating to dryness, melting the remaining mass, dissolving it in water, filtering and evaporating a second time.—Form, *flakey*.—Colour, *white*.—Taste, *pungent*.—*Deliquescent*.—Easily soluble both in *Water* and *Alcohol*.—Its acid decomposed in distillation.—Yields *Acetic Acid* on the addition of the *Sulphuric*.—Used in medicine as a *Diuretic*.

With *Ammonia* it produces, *Acetate of Ammonia* (*Liquor Ammoniacæ Acetatis P. L.*); prepared by saturating distilled *Acetic Acid* with *Ammonia*. — Form, *liquid*. — Taste, *urinous*. — Employed as a *diaphoretic*; sometimes also used externally as a *discutient*.

Order of attraction, *Barytes, Potash, Soda, Strontian, Lime, Ammonia, Magnesia, Metallic Oxyds, Water, Alcohol*.

Of Tartaric Acid.

Obtained from *Acidulous Tartrate of Potash* by means of *Lime*, or *Carbonate of Lime*, and the subsequent addition of *Sulphuric Acid*. May also be prepared by the *Sulphuric Acid* alone.

Crystallizes in small transparent scales or needles. Not altered by exposure to *Air*. Readily soluble in *Water*, and of an agreeably acid taste.

Easily decomposed by *Heat*.

Yields *Oxalic Acid* by treatment with the *Nitric*, and *Acetic* by digestion with *Water* and *Alcohol*.

Its more important saline compounds are,

we apply Sulphuric Acid to acetate of
tash of 4 parts of ^{the Sulph} Acid 13 Parts of acetate of
tash by Distilling this you obtain Acetic
id. it frequently contains a portion
Sulphurous Acid gas, which causes
a Decomposition of $\frac{1}{4}$ Sulphuric
id in $\frac{1}{4}$ Retort, of this it may be deprived
putting $\frac{1}{4}$ Acid into a clean retort
D with a little Black Oxide of
manganese, & a little Baryte. by this
you will get Pure Acetic Acid
Saturating Powder Subcarb. with Distil
ngan and evaporating it down to a Dry
ex.

The Pure State of Acetic Volatile & in
a Dry inflammable Nitric Acid,
mixes it into water and Carbonic Acid
it gives a little above the water

does not act on $\frac{1}{4}$ Simple Combustible
Substances but when combined with the
aromatic Benzene is by Acetic Acid
roughly $\frac{1}{4}$ on Point of Alcohol 2 Dr
Aromatic Camphire of 1 Dr
Lemon: 3, Oe. Pimento 3,

it is possible to obtain in the state of
Crystals but by being very volatile
it is difficult.

Sublimed tartaric acid is but in appearance so
of the tartaric as copper chloride. Under with all
the same, requiring only Acid Tincture
which is very common with Soda & by boiling
you will get a very salt the Acid of tartar
Tartaric Acid

in veg. acid, in a concrete form it is
radical Carbon & Hydrogen like that being
known as tartaric acid. When united with an excess of Potash
it forms $\frac{1}{2}$ Cream of Tartar (or Acidulous
Tartarate of Potash) which is sparingly soluble in water. The Impure state of it
is purified by solution to making white
Cream of Tartar.

It is met with in $\frac{1}{2}$ juice of grapes Apples &
(Acidulous Tartarate of Potash). We obtain the
Tartaric Acid from common Cream of Tartar
which changes to tartaric powder showing
except in acid this is dissolved in hot water then
mixing up a quantity of Carbonate of
lime forms with tartaric acid an
insoluble compound for many Tartarate of
lime, from which $\frac{1}{2}$ Tartaric Acid is formed
mixing up a quantity of Acid Sulphur equal
the weight of the chalk is used. The acid and
lime form Sulphate of lime
or Plaster & the Tartaric Acid is left.

Crystals of Tartaric Acid is very soluble
in water. Tartaric Acid is Dissolved
affording Crystals when poured into
water: but it is not so in Soda.
acid dissolves Argol of near hence
and by Calces Printures become
very common figures.

forms simple salts with most
the Oxides of Metals and also with
tash... The Rochelle Salt is an instan
when mixed with a salifiable basis
once obtained by adding to a solution
of Tartar Soda it is made
a union of these two preparations
is a mild Cathartic & tastes rather
than; in most of its properties
the Tartrate of Potash. —

Tartaric Acid is formed from wine
its acid and Argol when from which
White Argol

Took a Boiling Solution of Cream of Tartar
from under a Boiling Solution. Subsequently
16 1/2

1st. *Acidulous Tartrate of Potash*, prepared from *crude Tartar* (the spontaneous deposit from new wine) by solution and evaporation (*Potassæ super tartras* P. L.), or by crystallization under cooling.—Form *crystalline*.—Taste agreeably *acid*.—Of little solubility in *Water*.—Decomposed by the application of *Heat*; its chief products, an *Acid liquor*, and *Carbonate of Potash*, which has therefore been called *Salt of Tartar*.—Of various application in the arts, as in *Tinning*, *Dying*, *Hat-making*; employed also extensively in *Medicine*.

2d. *Tartrate of Potash* (*Potassæ Tartras* P. L.); prepared from the solution of the former, either by precipitating its excess of acid by means of *Chalk*, or by saturating it with additional *Potash*—Form, *crystalline*.—Taste, *saltish-bitter*.—Attracts moisture from the atmosphere; also very soluble in *water*; hence its former name *Soluble Tartar*.—Decomposed by heat, nearly in the same way as the *Acidulous Tartrate of Potash*.—Employed only in medicine, as a *mild Cathartic*.

3d. *Tartrate of Potash and Soda* (*Soda Tartarizata* P. L.); obtained, by evaporation and crystallization, from the union of *Soda* with *Acidulous Tartrate of Potash*.—Form of its crys-

tals, *prismatic*.—Taste, *bitterish saline*.—In most of its properties, and in its application, analogous to the *Tartrate of Potash*.

Order of attraction of Tartarous Acid, *Lime, Barytes, Strontian, Magnesia, Potash, Soda, Ammonia, Alumine, Metallic Oxyds, Water, Alcohol*.

Use confined to *Medicine*.

Of Oxalic Acid.

So called from being contained in the *Oxalis Acetosella*, or *Wood Sorrel*.

Obtained from *Sugar, Malt, Galls*, and various other substances, as well animal as vegetable, by treating them with *Nitric Acid*.

Form, *concrete*. Crystallizes in transparent quadrangular *Needles*. Taste, very *Acid*.

Readily soluble in *Water*: soluble also, and without decomposition, in the *Sulphuric* and other *Acids, Spirit of Wine, Æther*, essential and expressed *Oils*.

Easily decomposed under the application of the higher degrees of *Heat*.

Forms peculiar compounds with the *Alkalies*, with most of the *Earths*, and with several of the *Metallic Oxyds*.

Oxalic Acid

This Acid is existing in common Wood
trunks & the garden many other plants growing
naturally. I believe it is made of
Carbon & a diff. proportion of Oxygen

To Make it Take 5 parts of Nitric Acid diluted
with an equal part of Water, Part of 4
of loaf sugar, it should be heated very
gradually and made under a chimney to
prevent escape of ^{an} Nitric Acid Gas. The sugar
loses $\frac{1}{2}$ of its Oxygen & forms $\frac{1}{2}$ Oxalic
Acid you now have a solution of Oxalic Acid
& the Crystals will be formed by Evaporation
it may also be obtained from Gum Arabic
by the same process with Nitric
Soluble in twice its weight of Cold
Water, and a equal weight of boiling Water

Oxalic Acid does not act on Gold Platinum &c. but most
usefully on Iron which it readily
dissolves, for this purpose by Calico Printers
or taking out from ~~the~~

Compounds of this Acid with a salifiable
base are termed Oxylates, it is capable of
acting with Potash in 12 & a proportion

of sources as it is called in the Superoxylate

It is capable of forming with Soda and
then makes Oxylate of Potash

The Oxylate of Ammonia is a very useful
Compound indeed. You have nothing to do
than to saturate down by Oxalic Acid, Ammonia
& evaporate the solution.
Ammonia like $\frac{2}{3}$ other Alkalies is capable of
taking

The Oxylate of Ammonia is a delicate test
for Lime, because the Oxalic Acid forms with
the Lime an insoluble Compound
which is precipitated. To know $\frac{5}{7}$
quantity of Lime present filter it and
put it into a Crucible & apply heat

Citric Acid

It is existing in Chevre's Lemon
Lime

This Acid like Oxalic for use with Lime
an insoluble compound.

For obtain, it from Lemon Juice
and Carbonate of Lime, pour them
on a filter wash it & gradually dry

Crystals don't deliquesce in ² of Air

acts more powerfully on oxide of Iron
than Oxalic Acid

which takes place, and subsequent *Crystallization*.

Forms small granular or needle-like *Crystals*, of a sour but not astringent *Taste*.

More soluble in *Spirit* than in *Water*.

Burns in the open fire, leaving behind a hard insoluble *Charcoal*.

In close vessels, yields an *acid Liquor* and a *whitish saline Sublimate*.

A similar salt procurable from *Galls*, by *simple Distillation*; this also called *Gallic Acid*.

Remarkable for the property of producing a black colour with *Iron* and its saline compounds; hence the formation of *Ink*.

Hydro-carbon the basis of both these forms of *Gallic Acid*, but united with different portions of *Oxygen*.

Of Phosphoric Acid.

Sources of this acid various. Usually procured from common *Phosphorus*, by deflagration; or by slow combustion, under long exposure to *Air*; or by treatment with the *Nitric Acid*.

then Ick has been obliterated & may
be removed by various means...
do it your pencil the first with
syringe of Tincture and then follow
with Sulphuric Acid

then Ick when mixed with milk
of show itself until it is heated by
fire. Sulphuric Acid will thus act
on Metallic solution of various substances
on this property, of your words with
solution of Sugar of Lead nothing will
appear, but your fear over it, Sulphur
and Hydrogen water.

Phosphoric Acid

To make it Take Nitric Acid placed in a Florence Air flask and put in Stick of Phosphorus under of Nitric fumes close & Grasp off —

It forms Compound with Salifiable basis called Phosphites

a solution of Phosphites is capable of precipitating Golden Solution

Strong Phosphoric Acid is produced when Phosphorus enters into rapid combustion

Phosphoric Acid may be concentrated by evaporating the water —

Phosphoric Acid has not that Corrosive effect on Animal Matter as Sulphuric Acid

Glaf of Phosphorus is rather dele-
rious when exposed to the
atmosphere...

It combines with the Salphur
Basis and forms Phosphates.
(Phosphoric Acid Dues)

Phosphate of Soda is a mild Cathartic
it is more pleasant than the Sulphate
of Soda or Sulphate of Magnesia



Differs in its sensible properties according to the degree of *Oxygenation* : hence distinguished into *Phosphoreous* and *Phosphoric Acid*.—In the latter form, generally liquid ; but by evaporation and cooling, crystallizable into quadrangular *Prisms*.

Taste, intensely acid, but not corrosive. *Specific gravity* 2. 687.

Has a strong attraction for *Moisture*.

Under the application of the higher degrees of *Heat*, fuses into a transparent *Glass*, capable of re-assuming its acid properties by long exposure to *Air* or maceration in *Water*.

With the *Alkalies*, *Earths*, and *Metals*, forms particular compounds, the chief of which is, *Phosphate of Soda* (*Soda Phosphorata* P. Edin.) ; obtained by uniting *Phosphoric Acid* and *Soda* to the point of saturation, evaporating, and cooling. Form *crystalline*.—Taste not unpleasantly saline.—*Efflorescent*.—Dissolves readily in *Water*.—Forms a triple salt with *Ammonia* (*Sal Microcosmicus*).—Lately introduced as a *thartic*.

May, like most of the other acids, be partially or totally decomposed, by treatment with different *inflammable substances*.

In distillation with *Charcoal*, re-produces *Phosphorus*; this therefore considered as its *Radical*.

Order of attraction, *Barytes*, *Strontian*, *Lime*, *Potash*, *Soda*, *Ammonia*, *Magnesia*, *Alumine*, *Metallic Oxyds*, *Water*, *Alcohol*.

Of Prussic Acid.

Obtained from *Prussiate of Iron* (*Prussian Blue*), by boiling it in a solution of *Potash*, adding *Sulphuric Acid* to the liquor when filtered, and submitting it to distillation; or by distilling a mixture of *Prussiate of Potash* and diluted *Sulphuric Acid*, and re-distilling the product from *Carbonate of Lime*.

Form, *gaseous*.

Unites readily with *Water*.

Combines with *Alkalies*, *Earths*, and *Metallic Oxyds*; but these compounds easily decomposed by *Carbonic Acid*.

Ferruginous Prussiates—Their uses.—With *Oxyd of Iron* re-produces *Prussian Blue*.

Consists of the same elements as *Carbonate of Ammonia*, but in different proportions.

Order of attraction, *Barytes*, *Strontian*, *Potash*, *Soda*, *Lime*, *Magnesia*, *Ammonia*, *Metallic Oxyds*, *Water*, *Alcohol*.

Prussic Acid

May be Made Gaseous.

Prussic Acid ought not to be exposed to the Atmosphere & light or the Prussic acid will lose its nature.

It is principally of Animal Origin or Substance which contain Azote. May be prepared by mixing Dried Ox blood with Subcarbonate of Potash (Blood 3ij Alkali ^{Pot. Subcarbonate} 3i) then heated till they no longer puff up; it must be exposed to a red heat until it runs to flame. This substance thrown to water produces the Prussiate of Potash which is very liable to decomposition.

Prussian Blue consists of equal parts of Oxide of Iron & Prussic Acid. It is rather inflammable. Detonates when mixed with Oxymuriate of Potash, when struck yields an explosion Hyd. Ammon. Gas & Carb. Ammon.

Blow alone will yield Prussic Acid
by Distillation & Spirit of Hartshorn in Leaves of
Green Kernels of Apricot
To School we are indebted for the Compara-
tion of this Acid.

Take 3 table Spoonful of Charcoal powder
the same quantity of Sublimed. of Potash
Subjecting them 2 substances to heat & then
put in Mixture of Ammonia & Decaying
or 2 more will take place

There is great tendency in Prussic Acid
to Decompose and form Carbonate of
Ammonia

When concentrated this acid is most violently poisonous
Prussic Acid is very Volatile - and stands lowest
as to the Salifiable basis of all Acids.

We obtain it in Liquid form because Water
Absorbs $\frac{1}{4}$ Gas so readily, but this should not
be exposed to Atmospheric Air

It will boil at $\frac{1}{2}$ temperature of 85 or 90 of Fahrenheit

Original Bern 3y and 4y of Mercury leave them together
& you get a Solution of Prussic Acid of Mercury

a Solution of Prussiate of Mercury & Iron filings
put into a retort and a little concentrated
Sulphuric Acid; it will cause a Decomposition
& by it you obtain Prussic Acid - in a Solution

of all the acids & the 3 acids will be lost & you will
obtain the gasous Prussic Acid - which may be absorbed

In order to act on Phosphate it must
be combined with Alkali:—

Nitrogen Gas by the French Chemists, when combined ^{with}
hydrogen called Hydro nitrogen Acid

A green Sulphate of Iron is precipitated
by Purpuric Acid but becomes
blue when exposed to the Air,

Purpuric Acid. Alkalies & earths seem
to have the strongest affinity for
it.

The Purple Salts formed by this Acid &
alkaline earths, and carbonic acid
are very unlikely to Decompose

Purpuric Acid is capable of 2 States of Ox-
idation first the Purpurous Acid
2nd by the Purpuric Acid

A solution of Purpuric Acid will precipitate
Metals but not the Earths therefore valuable
vs. a solution of Copper it precipitates
a brown color: a solution of Nitrate of Lead
will precipitate a white color, a solution
of Nitrate of Silver.. it precipitates a yellow.

For example Purpuric Acid & Purpurous Acid the solution of Potash
at then together the color will be changed to brown. If of other
appear the color it will be brown, after then pour the
whole upon a filter the color will be brown, gently to dryness
at Purpuric Acid.

To Make the Salt.

Take caustic Potash add to it powdered Prussian
Blue, which changes its color to a brown
and continues adding the P^r Blue until
the color of it is no longer changed
then filter it and you obtain a clear
solution of ferrugineous Prussiate of Pot.
which is much more permanent
the Sulphur which it is very liable to contain
is got rid off by -- By evaporating this
solution you obtain the Prussiate of
Potash in crystals. The Mass that is
left on the filter will you find contain
a portion of Prussian Blue

Benzoic Acid called Soluble in water
when in a solid form from Gum Benzoin made by
heating it, found in Balsam of Peru & Japan.

It has been discovered combined with Lemon
and the Urine of Horses & Cows

24 Parts of hot Water will dissolve one part of
it. Alcohol dissolves it completely

Powder of Benzoin subject it to heat and
cause the Vapor by a Drum on which
on the sides the Acid adheres. To Purify
it subject it to a strong pump and again sublime
it, when beautiful white crystals will
be obtained. From: Another way that the Succinic
Acid is obtained an expensive Acid. Succinate of Am
is $\frac{1}{2}$ only useful form of it?

OF EARTHS.

Though comparatively *few* in number, met with under an infinite diversity of appearance, and in such abundance as to constitute the basis of the more solid parts of the globe.

Seldom if ever found free from foreign admixture: when purified artificially, distinguished by their want of *Tenacity*, *Fixity*, sparing solubility in *Water*, *Insipidity*, want of *Odour*, incapacity of communicating a tinge to *Glass*, and their specific gravity not exceeding that of *Water* more than in the proportion of five to one.

These characters possessed more perfectly by some of the earths than by others: hence their distinction into *saline* and *insipid*.

All the earths soluble in one or other of the *Acids*: cannot however be precipitated from their solutions, like the *Metals*, by *Prussiate* of *Potash* or of *Lime*.

Infusible even by the most intense degrees of *heat*, unless in a state of mixture; viz. with *each other*, with *Alkalies* or *other Salts*, or with *Metallic Oxyds*.

The earths at present known are, *Barytes*, *Strontian*, *Lime*, *Magnesia*, *Alumina*, *Silica*, *Zirconia*, or *Jargon Earth*, *Glucina*, and *Yttria*.

Of these *Lime*, *Alumina*, *Silica*, and *Magnesia*, by far the most abundant and useful.

Of Barytes, or Ponderous Earth.

Found in combination, 1st. with Carbonic Acid, *Carbonate of Baryt*; 2d. with Sulphuric Acid (*Sulphate of Barytes*, *Baroselenite*;) or 3d. with Sulphuric Acid, *Silex*, *Sulphate of Alumina*, *Sulphate of Lime*, and *Petroleum*, (*Liver Stone*.)

Obtained in a caustic or separate state, by exposing a mixture of *Carbonate of Barytes* and *Charcoal*, or *Nitrate of Barytes*, to a strong heat.

Colour greyish.—Taste caustic, poisonous.

Discovered by Davy to be a *Metallic peroxide*.
—*Barium*.—Means of obtaining, and properties.

Slakes like *Lime* on exposure to air, and imbibes water with avidity, forming with it a powerful cement.

Malic Acid *Wiley's Chemical Dictionary*

useful in forming with Lime a
soluble compound. so by means of
this acid you obtain Lime from Barytes
This acid exists in a variety of fruits. -
In drying the Malic Acid forms
other acids by Alcohol. Malic Acid is
not capable of Chrysallization.
Suberic Acid

may be obtained by applying heat
Pieces of Cork -

1. Powder and distill with water
1 lb of cork applying heat, filter & evaporate
dryness. Formic Acid Lactic Lactic
acid. &c

~~Properties~~

1. It is a weak, pungent, of heavy
dense color of brown. It is very volatile
when

it has the property of boiling on a large scale
water, which would not be the case if it
were the case. It is very easily heated to
boil. It is very easily formed from the acid
I

The first of these, being the title of the solution as
given by the author, is not a solution of the problem
as stated with regard to the first part of the
problem, but is a solution of the second part.
The second of these, being the title of the solution as
given by the author, is not a solution of the problem
as stated with regard to the first part of the
problem, but is a solution of the second part.
The third of these, being the title of the solution as
given by the author, is not a solution of the problem
as stated with regard to the first part of the
problem, but is a solution of the second part.

The fourth of these, being the title of the solution as
given by the author, is not a solution of the problem
as stated with regard to the first part of the
problem, but is a solution of the second part.
The fifth of these, being the title of the solution as
given by the author, is not a solution of the problem
as stated with regard to the first part of the
problem, but is a solution of the second part.

Soluble in about 20 times its weight of cold, and in less than twice its weight of boiling water (*Barytic Water*,) from which in cooling it crystallizes in transparent prisms.

Imparts a lemon colour to the flame of *Alcohol*.

Unites with the *Sulphuric Acid* into a compound requiring for its solution 40,000 times its weight of *Water*.

With the *Nitric* and *Muriatic Acids*, forms crystallizable salts.

Unites with *Sulphur* into a species of Hepar (*Sulphuret of Barytes*.)

Order of attraction in the moist way, *Sulphuric Acid*, *Oxalic*, *Succinic*, *Fluoric*, *Phosphoric*, *Saccho-lactic*, *Nitric*, *Muriatic*, *Citric*, *Tartaric*, *Arsenic*, *Lactic*, *Benzoic*, *Acetic*, *Boracic*, *Sulphureous*, *Carbonic*, and *Prussic*; *Sulphur*, *Phosphorous Water*, *Unctuous Oils*; in the dry way, *Phosphoric*, *Boracic*, *Arsenic*, *Sulphuric*, *Succinic*, *Fluoric*, *Nitric*, *Muriatic*, &c.; *Potash*, *Soda*, *Sulphur*, *Oxyd of Lead*.

The *Carbonate of Baryt* and other forms of this earth, remarkable for their violent effects, when exhibited internally.

Used principally as a test for ascertaining the presence of *Sulphuric Acid*.

Of Strontian.

Found either in combination, with carbonic acid, *Strontianite*; or with sulphuric acid, *Sulphate of Strontian*.

Best obtained in a separate or pure state from *Carbonate* or *Nitrate* of *Strontian*, by exposing them to a strong heat, as in the preparation of *Baryt*, to which it is analagous in many of its properties.

Discovered by Davy to be a *Metallic Peroxide*—*Strontium*.—*Strontian* nearly resembles *Barytes*—Soluble in 160 parts cold water, in less of hot water—The solution on being evaporated affords crystals of pure *Strontian*.

Remarkable for the brilliancy of the flame which it exhibits when treated on charcoal by the blow-pipe; also for the red colour which it imparts to the flame of *Alcohol*.

The order of its attraction nearly the same with that of *Baryt*, though inferior to it in degree.

Of Lime or Calcareous Earth.

Found abundantly in different parts of the world, combined with *Carbonic Acid* in the forms

Stuntian

little is about 100 times the weight of the

Stuntian - my experience from the

history of the Stuntian

Stuntian is a very common

Stuntian is a very common

Stuntian is a very common

Stuntian is a very common

Stuntian is a very common

Lime sometimes found with clay when it is called
Marl

The Composition of Mortar must be varied according to the purpose intended for. Sand & Lime is $\frac{1}{2}$ Bar of all kinds of Mortar

These stones & bones are everywhere to be seen & are
these stones & bones, Siliceous & lim. must be very easily.

refining & Sugar Making off the River from

of *Chalk*, *Lime-stone* and *Marble*—also in the bones and horns of animals, and the shells of fishes and the eggs of birds—Prepared for various purposes from *Carbonate of Lime*, by the continued application of a strong heat.

Form *concrete* or *powdery*. Taste *hot, pungent, caustic*.—According to Davy, is a *metallic peroxide*.

Soluble in about 500 times its weight of water (*Liquor Calcis* P. L.) Changes vegetable blue colours to green.

On being suddenly moistened, emits both *Heat* and *Light*, losing at the same time its attraction of cohesion, *Slaked Lime*. The same takes place spontaneously on exposure to *Air*. On further exposure attracts *Carbonic Acid* from the *Atmosphere*.

Though infusible *per se*, promotes very powerfully the fusion of most of the other earthy Bodies: hence its use in working metallic ores, more especially those of *Iron*.

Melts with *Borax* and *Microcosmic Salt*, without effervescence. Melts also with *Oxyd of Lead*.

Use of *Lime* as a cement—composition of mortar—substances which give it the property of resisting the action of water—importance of

Lime as a manure—methods of analysing Lime Stones.

Sulphate of Lime, called also *Gypsum* or *Senenite*, soluble in about 500 parts of water, but in less if the water contain an excess of acid—when heated to redness falls into a soft white powder.

Plaster of Paris—its uses.

With *Nitric* and *Muriatic Acids* forms *deliquescent Salts*.

Phosphate of Lime obtained from burnt bones, sometimes found native, as in *Apatite*.

Fluate of Lime found in Derbyshire, Cornwall, and various parts of the world—When heated *phosphorescent*—With *Sulphur* forms a calcareous *Hepar*; with *Phosphorus* a liver-coloured compound, which yields *Phosphureted Hydrogen Gas* on the affusion of water; with *unctuous substances*, peculiar *Soaps*.

Order of attraction in the moist way, *Oxalic Acid*, *Sulphuric*, *Tartaric*, *Succinic*, *Phosphoric*, *Saccho-lactic*, *Nitric*, *Muriatic*, *Fluoric*, *Arsenic*, *Lactic*, *Citric*, *Benzoic*, *Acetic*, *Boracic*, *Sulphureous*, *Carbonic*, and *Prussic*, *Sulphur*, *Phosphorous*, *Unctuous Oils*, and *Water*; in the dry way, *Phosphoric*, *Boracic*, *Arsenic*, *Sulphuric*, *Succinic*, *Nitric*, *Muriatic*, *Fluoric*, *Lactic*, *Ben-*

is Quick Lime that is used for Making
mortar. The more labor used in Making
the better. as it more effectually fills
the interstices. When Mortar is kept
a long time excluded from $\frac{1}{2}$ Air, it be-
comes more durable. The Setting of
mortar depends on 2 or 3 causes. If it
to be exposed to water it should be well
before it is exposed to it. It should be
back of some space. It should be well incorporated
and Water Cement is also made with
real parts of Abertown as I found.

is often employed in Agriculture, for this it
should be free from other matters. Besides removing
the effect of a cold earth, it hastens putrefaction
by this means is useful in Vegetation. It
should be used only on Land where there is a
large quantity of Vegetable Matter, but not
otherwise, as it would injure the Land very much.
Lime Stones which contain Magnesia
The varieties of Marble consist of Carbonic
acid & Lime.

Many engineers say that the ancient Roman
engineers that Marston should be used to
engineers and many have used it from
the Atlantic.

The use of iron in forming cement is used in
foundations when the shaft of an anchor is broken
when the stone is used in water to form
a strong foundation.

From one method has the property of giving
hardness to cement under water.

The iron then is off water from by Blast
when hammering is used to produce a
mix with water from a strong cement.

To fish ponds take 2 parts of Lime containing
1 part of Portland & 3 parts of ⁴ ^{thick} clean sand
forms a most excellent water cement.

out of blind matter or made by 2 measures
barkham. Linn. Hume. Linn.
the minerals and fossils with the minerals
Linn. with the minerals and fossils with the minerals

Sulphate of Lime is useful in
agriculture. Sulphate of Iron such
unify calcareous earths.

To obtain Phosphate of Lime you will
wash bone ash, then add Phosphoric Acid
it, and then precipitate the pure Phos-
phate of Lime by Ammonia.

Strong Lime is made as very big
measures of sifted Lime & of sand & of
and Ashes & Bone meal & after some time
it is added

even much water in Stone nearly finished
specimens in that are arranged fully
in the calcareous parts of Lime
phosphate of Lime. Slightly soluble in water & gives
simple clear taste. It may be precipitated
highly red by Alcohol but it is light to be very
in the contents in State of Limestone or large quantity
of water of charged with it. Carbon may be
made very common with the sulphate of Lime
the Carbon of Stone. The lime may be
very easily

Magnesia

When Pure a white Powder, sparingly soluble
in water but like Lime can absorb a portion
and appear Dry. Alkaline properties slight, like
impossible in its greatest heat. It will combine
with Sulphur only. (of the Lanthan tribe)
Can may suggest from being an absorbing compound
that its Carbonic Acid is driven off
The water contains it in considerable portions
Form a solution of Sulphate of Magnesia
The Magnesia used in Medicine is prepared
It is precipitated by ^{the} Carbonate of Soda. Or
The Patent Magnesia is made exactly by the same
process. The precipitation when Dry is cut
into square pieces by means of a Saw.
Magnesia is made by burning the Carb.
of Magnesia in a Crucible

^{acid}
Sulphuric entirely dissolves Sulphuric
Acid, but it will not Sulphate of Lime
Oxide of Ammonia will precipitate Nitrogen
of Lime but not Magnesia therefore a test for
Lime. Pure Ammonia will throw down Mag-
nesia but not Lime

zoic, and *Acetic Acid*; *Potash*, *Sulphur*, *Oxyd of Lead*.

Used in *Dying*, *Bleaching*, *Tanning*, *Sugar-baking*, and various other arts, besides its application in *Medicine*.

Specimens of Stones in which *Lime* forms a principal part.

Of Magnesia.

Supposed by Davy to be a *Metallic Peroxide*.

Prepared from a solution of *Sulphate of Magnesia*, by the addition of *Carbonate of Potash*, and subsequent exposure of the washed earthy precipitate to a strong and continued heat (*Magnesia P. L.*)

Form *pulverulent*. Colour *pure white*. Taste *insipid*.

Requires for its solution 7,900 times its weight of water. Tinges vegetable blues of a light green.

Infusible without addition, even in the most intense degrees of heat, by which it is merely contracted in its dimensions; but melts into a glass with *Lime*, *Microcosmic salt*, or *Borax*, or with a mixture of *alumina* and *silica*.

Unites with all the Acids. With the Sulphuric Acid regenerates Sulphate of Magnesia (*Magnesiæ Sulphas P.L.*) With Carbonic Acid, Carbonate of Magnesia (*Magnesiæ Carbonas P. L.*)

May be combined, in small proportions, with Sulphur.

Order of attraction in the moist way, Oxalic Acid, Phosphoric, Sulphuric, Fluoric, Arsenic, Saccho-lactic, Succinic, Nitric, Muriatic, Tartaric, Citric, Lactic, Benzoic, Acetic, Boracic, Sulphureous, Carbonic, and Prussic, Sulphur, Phosphorus, and Water; in the dry way, Phosphoric, Boracic, Arsenic, Sulphuric, Fluoric, Succinic, Nitric, Muriatic, Lactic, Benzoic and Acetic Acid; Potash, Sulphur, Oxyd of Lead.

In common use in disorders of the *Primæ Viæ* as an *antacid* and *laxative*, and in the form of steatite as an ingredient in the Manufacture of the finer kinds of Pottery.

Specimens of Stones to which it gives a distinguishing character.

Of Alumina.

Obtained in its purest form from a solution of common Alum, or Sulphate of Alumina, by the

Magnesian contents & composition of
various stones. In Steatite or Soap Rock
contents. The Lapis Albarius it is in (or
not stone) -

The Rock called Temperstone is a Magnesian Specimen
The Heckel & Polston are Magnesian Specimens.

Argil

Argil is obtained from Common Alum
the commonest rock 3 is found by the sea on ~~the~~ ^{the} ~~left~~ ^{right} ~~side~~ ^{side}

Argil is found in Argil, its deposit to Chertall is
the commonest ~~rock~~ ^{stone} ~~is~~ ^{is} ~~found~~ ^{found} ~~in~~ ⁱⁿ ~~the~~ ^{the} ~~country~~ ^{country}

The commonest fossils of Argil are Chertall
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engine has a strong attraction for ~~vegetable~~ ^{animal} matter
as compared to the Dryden one - more so

aluminum from as frequently as compressed liquid air
in which is present to the degree

is many be got rid of by Boiling in K. Shaving it under
this divides the particles very much & the Sulphating Pro-
cess be washed out. For Woolen this Ferri Sulph: doubt signifi-
cantly does harm when for Silk the
or Solution of Chlorine be be used with a solution
argue The coloring matter of the Acum. is soluble
in Cassia with it & forms the coloring matter
and dark Vegetable Matter answers equally well as Nadder

much Chalk in Magnesian Specimen. - See Sketch.

large in many places and making Bunka. but it
by becoming more dense in appearance of becoming black
containing black

in which may be made between the House and the Senate. The House has the power to impeach the President and the Senate to try him. The House also has the power to impeach the Judges of the Supreme Court and the Judges of the inferior Courts. The Senate has the power to impeach the Judges of the Supreme Court and the Judges of the inferior Courts. The House has the power to impeach the Members of the Senate and the Senate has the power to impeach the Members of the House. The House has the power to impeach the Members of the Executive Branch and the Senate has the power to impeach the Members of the Executive Branch. The House has the power to impeach the Members of the Judiciary Branch and the Senate has the power to impeach the Members of the Judiciary Branch. The House has the power to impeach the Members of the Legislative Branch and the Senate has the power to impeach the Members of the Legislative Branch. The House has the power to impeach the Members of the Executive Branch and the Senate has the power to impeach the Members of the Executive Branch. The House has the power to impeach the Members of the Judiciary Branch and the Senate has the power to impeach the Members of the Judiciary Branch. The House has the power to impeach the Members of the Legislative Branch and the Senate has the power to impeach the Members of the Legislative Branch.

Thais soluble in Water it may be suspended in
When Dry it will absorb a large quantity
of Water.

3 Argie 2 Lime, Magnesia when united will form a
Heat

Argie has no effect on Argil. This is not correctly
ascertained

Argil has a strong attraction for Vegetable &
animal Matter, which is very easily
removed by Nature.

But above all it is useful in Dyeing, as a
Mordant

Fullers earth Consists of argie & Lime. therefore when applied
to Vegetable or animal Matter (which it is employed by some
for) on account of its strong attraction for them it removes
them

Argie has a particular action on $\frac{2}{3}$ Metallic
Alyds. forming Ochre or what was formerly called
Boley. They are all ⁱⁿ fused in (cannot be done)

Clay contracts a great deal when a high heat
is applied therefore in making Pottery it is necessary
to add a certain quantity of Silicious Matter
in to prevent its cracking

The Russian black flint is the best for blackening, owing
to its Carbonaceous Matter, it is ground to a very fine
powder before it is used for making Pottery. This is
the common Tobacco Pipe Clay makes the common
Pottery. They used to sift the clay but it is injured the
most necessary much. The felspar being very insoluble &
expensive than lime. The felspar is mixed with the
clay by adding for the workman at a furnace for
firing make an oven about 100 $\frac{1}{2}$ Pounds.

day is not so readily acted upon by the strong acids as
Sulph. Min. Nitre &c
Fixed Alkali readily decomposes therefore exceedingly useful
in analysis as for example when with Lime & Magnesia
which it will not decompose.

by the addition of Potash, Soda, or Ammonia; hence
its present denomination.

by Alkali Form powdery. Colour, when dry, pure white.
Feel unctuous. Taste insipid. Smell, when
breathed on, earthy.

Insoluble in Water. When moistened be-
comes plastic, and contracts and hardens when
exposed to the higher degrees of Heat; therefore
the basis of the different kinds of Pottery. After
baking is no longer capable of becoming plastic.

Soluble in the humid way by Alkalies.

Fusible, with effervescence, both with Micro-
cosmic Salt, and Borax, fusible also with Lime.

Combines with most Acids, though with dif-
ficulty, except under precipitation, and pro-
duces with them compounds which are more or
less astringent. The most important of these,
the Sulphate of Alumina or common Alum, pre-
pared from the decomposition of Argillaceous
Schistus. This properly a triple, sometimes a
quadruple, salt. Form of its crystals octohedral.
Taste astringent. Soluble in about 14 times its
weight of cold, and somewhat more than an
equal weight of boiling water. Undergoes wa-
tery fusion, and parts with its water of crystal-
lization, on exposure to Heat (*Alumen exsicca-*
tum P. L.) When calcined with certain in-

They now strain in mixed with Water thus a Scina.

Plants, which stand & dry up & become brittle & they are melted & Porcelain is made the most of a Concretion of common earth melting at a heat 58 broken is looking like glass or something the appearance of wax. Common earthen ware looks granular

flammable substances, as yolk of egg, or sugar, produces a compound, which takes fire spontaneously on exposure to Air (Pyrophorus of Homberg.)

Has, of all the Earths, the greatest attraction for Metallic Oxyds; has also a strong attraction both for Silica and Magnesia.

Order of attraction of Alumina in the moist way, Sulphuric Acid, Nitric, Muriatic, Oxalic, Arsenic, Fluoric, Tartaric, Succinic, Saccho-lactic, Citric, Phosphoric, Lactic, Benzoic, Acetic, Boracic, Sulphureous, Carbonic, and Prussic; in the dry way, Phosphoric, Boracic, Arsenic, Sulphuric, Nitric, Muriatic, Fluoric, Succinic, Lactic, Benzoic, and Acetic Acid; Potash, Sulphur, Oxyd of Lead.

Uses, comprehending those of Sulphate of Alumina, Dying, Tanning, Printing, Silvering, Painting, Pottery, Medicine, &c.

Specimens of Stones containing this Earth in considerable proportion.

Of Silica.

Exists, with very little mixture, in Sand, Gravel, Flint, and Rock Crystal.

Porcelain is made the most fusible kind of Glaze as Borax. The earth of Porcelain for 72 earthen. They form the Chinese Porcelain or Japan ware.

The Glaze of common ware is made of Oxyd of Lead 5 lb & 1/2 of Soda. Mix them with one 4 lb of Soda. Fire in a dry way. The Glaze of fine ware is made of Soda 10 to 20 lb & 1/2 of Soda. Fire in a dry way. The Glaze of fine ware is made of Soda 10 to 20 lb & 1/2 of Soda. Fire in a dry way.

Porcelain is made that it strikes fire with steel & does more the strongest than with impunity

Silica is the most abundant substance in nature. It is the basis of all rocks & minerals. It is also the main component of glass & porcelain. The attraction of Alumina for metallic oxyds is very strong. The attraction of Alumina for silica & magnesia is also very strong. The order of attraction of Alumina in the moist way is as follows: Sulphuric Acid, Nitric, Muriatic, Oxalic, Arsenic, Fluoric, Tartaric, Succinic, Saccho-lactic, Citric, Phosphoric, Lactic, Benzoic, Acetic, Boracic, Sulphureous, Carbonic, and Prussic. In the dry way, the order is: Phosphoric, Boracic, Arsenic, Sulphuric, Nitric, Muriatic, Fluoric, Succinic, Lactic, Benzoic, and Acetic Acid. Potash, Sulphur, and Oxyd of Lead are also attracted. The uses of Alumina are many, including dyeing, tanning, printing, silvering, painting, pottery, and medicine. Specimens of stones containing Alumina in considerable proportion are found in various parts of the world. Silica is found in sand, gravel, flint, and rock crystal. Porcelain is made from a mixture of silica and alumina. The most fusible kind of glaze is made of silica and borax. The earth of porcelain is made of silica and alumina. The Chinese porcelain and Japan ware are made of silica and alumina. Porcelain is made that it strikes fire with steel & does more the strongest than with impunity.

in the last has been taken they are seen by 1 after an
in demand as a common demand on ground
the are even to be 1, as to branding them they are
tensands furnished by other workmen in Germany
Polish & American They are heated again
under a fire in a furnace where they take Brandy as
common Paper them in an 1/2 degree and then
by degrees that are been the property of making
element to be used for paper
the 7 Copper Green Cobalt Blue, Gold
Silver & Yellow

is ground up the Apperholdden and printed them as
from plates. Another they are afterwards printed
by the 1/2 degree. The common and some are
used with the 1/2 degree of heat.

Silica or Silica

Flint contains a large portion of it & many other
stones

It can't be Crystallized by art but under peculiar
circumstances,

It is capable of absorbing $\frac{1}{4}$ of its weight of water
is infusible by itself in the strongest heat
Nitric acid acts on it very strongly

Silix has a very strong attraction for $\frac{2}{7}$ Alkalies.

If 1 part of powdered Silix & 3 Parts of Carb: of Soda, heat them together and it will afterwards dissolve in Water. Then by pouring in Acid: Muriat: you precipitate the earth, Silix, from the Acid having a stronger attraction for the Alkali than the Alkali has for $\frac{2}{7}$ Silix.

Silix is not acted on by Oxygen and any Combustible Substances.

Pure caustic Barytes seems to act on it in the same Manner as $\frac{2}{7}$ Alkalies.

Silix is Obtained in a State of Purity from $\frac{2}{7}$

When Silix is melted with a large portion of Alkali it forms a Glass & Salts in Water but has combined with a smaller proportion of Alkali it forms Glaze. This Substance was known a very early period in British Museum in form of beads which they used to give as ornaments to their vessels.

It was not till the 3rd century that glass
came in general use for ornaments
and then in enormous quantities
part Silica & 2 Parts of calcined Borax
the form a fine glass, this is the
improved made to imitate various
colours. They are colored by various Metals.

Alkali used in Britain for Making Glass
Sea Sand this is prepared by washing best
brown Shagbark Wood
and for common purposes they obtain
the Alkali from Wood Ashes & Kelp.

Green tinged glass is owing to the pre-
sence of Iron: a portion of Arsenic
is used to prevent the glass from backing
when exposed to variety in temperature
It also causes the work to be in

Parts of Silica with one quarter of potash & alkali
is not less than 7 over Parts.

It is used in Making the best Glass

It is very useful in Making Glass
for Optical Purposes

The glass used in the is an only
known to be in it.

Borax

Flint glass is acted on powerfully by Sulphuric
Hydrogen

in China.

White oxide of Arsenic is a powerful flux. It is
in every way a better flux than arsenic
it is in excess it becomes soft & dangerous
it is unsafe for drinking vessels.

It is used for making some of the
colored glasses.

When Purple Tungsten is heated
the whole with a blackish color

Silica mixed with twice its weight of Soda
forms a glass but the soluble in water
is Deliquescent. Silica with the same quantity
of alkali is melted also, a Perfect glass can
be formed when the alkali is $\frac{1}{2}$ the mass
of the Silica

May be separated from most of these by fusion with either of the fixed alkalies, and re-digestion in the nitric or any other of the stronger acids.

Colour *white*. *Inspid*. *Dry* to the touch. Sparingly if at all soluble in *Water*. Specific gravity 2.66. *Infusible*.

Soluble in the *Fluoric*, but in no other acid.

Soluble also in the solution of either of the *fixed Alkalies*, by the assistance of heat. Precipitates and combines with lime on being added to lime water; unites too in the humid way with some of the metallic oxyds.

May be fused with *Lime*, *Microcosmic Salt*, or *Borax*; but much more readily with *Potash* or *Soda*: hence the preparation of the different kinds of *Glass*.

When melted with a large proportion of *Alkali*, forms a deliquescent compound, (*Liquor Silicum*,) the decomposition of which by an *Acid*, furnishes a gelatinous precipitate, thought to be soluble in about 1000 parts of *Water*.

Order of attraction in the moist way, *Fluoric Acid*, *Potash*; in the dry way, *Potash*, *Boracic Acid*, *Phosphoric Acid*, *Oxyd of Lead*.

Used principally in *Polishing*, *Painting*, and *Glass-making*.

Specimens of Stones of which this Earth forms a principal part.

Of Zirconia, or Jargon Earth.

Combined with silex, iron, and nickel in the stone called *Jargon* or *Zircon*: found also in the *Hyacinth*.

Obtained from these by fusion with potash, and subsequent solution in, and precipitation from the *Muriatic Acid*.

Colour, *white*. Insoluble in water; when moist, *semi-transparent*. Specific gravity estimated at 4.300

When heated in contact with charcoal, is imperfectly vitrified, and becomes of sufficient hardness to strike fire with steel.

Resembles *Silica* in its action on *Metallic Oxides*, and *Alumina* in forming astringent compounds with the *Acids*: but differs from both in being insoluble in the *Fixed Alkalies*.

Order of its attraction as yet unknown.

Of Glucina.

Obtained by fusing *Aquamarine*, the *Beryl*, or the *Emerald* with three times their weight

the alkali is volatile it is driven off by the
heat of the solar rays & keeps behind therefore
by using your stove you remove nothing there is only
a loss of fuel to drive off the alkali

Acacia black, brown & Shagreen being
1 lb 30 Parts 16 Part, Pearl ash or Soda
ash 2 of Charcoal. This is brought
a red heat & is used for a purgative
also. Glass glass is used for Diamond &
softer than common glass

10 P of wh. Sand 33 Pearl ash
Oxide of Lead 11 Part of Nitre & about
200 parts weight of the whole of the material
in this glass. Phosphorus is made by drawing
it in a nail.

Glass glass is made as they
10 Part Sand 10 of Sub. Soda 10 P of
manganese is a kind of

Powder of Borax may be useful these
materials are given 30 hours under a heat
from the furnace with mounds

in common glass is made of common
materials

Materials are melted in a large
cup or glass

When the materials are put into a vessel
they are all heated to a point that they are

Tracking after their Prongs they now take them
together for 30 hours or thereabouts.

The Mathematical Hammer - From the
prongs is over the next is the Defensive
Prongs.

After the glass is properly formed it is
by

from principal instrument as a ^{Simply} ~~Prong~~
Globe

The glass is suddenly cooled after it has
made in with rapidly with. Therefore
after it is made with the glass they see
it. Sent to the anatomical furnace
which comes is gradually.

30000 from the
present from the
order to purchase the land
and they it with a view to
right of the land the whole of the
them for a few years
man is not satisfied by the
action but it does the
than the law is right to be
before the law and since the
man from which the land may

Glucina

Causes Glucina because the earth forms
from it in the same manner as Potash

Itoria

It is called Glucina in former times
and is found in the same manner as Potash
in the same manner as Potash

As a very rare earth

Remarks on the Glucina

The Glucina is found in the same manner as Potash
in the same manner as Potash

Causes & Itoria form a great part of the
of the Glucina. If the Glucina is removed
as Itoria & Glucina is the same because
it is the same.

Water is necessary to the Glucina for the
of the Glucina.

of *Potash*, separating the *Silica* and *Alumina* by the usual methods—then adding to the solution *liquid Carbonate of Ammonia* in excess, boil until the *Glucina* precipitates.

Colour, *white*. *Insipid*. *Insoluble* in water. *Adhesive* to the tongue. *Infusible per se*; but melts with *Borax* into a transparent *Glass*.

Soluble in most of the *Acids*, and likewise in the solutions of the fixed *Alkalies* and of *Carbonate of Ammonia*.

Its *saline* compounds slightly *astringent*, and of a *sweet taste*: hence its name.

Its affinity for *Acids* intermediate between that of *Magnesia* and *Alumina*.

Yttria.

Found in the *Gadolinite*, a Swedish stone, so called from this earth being discovered in it by *M. Gadolin*.

Procured by treating the powdered mineral with *Nitric* and *Muriatic Acids*—evaporating the solution to dryness—diluting with *Water* and precipitating the *Yttria* with *Ammonia*.

Colour of this earth *pure white*—Without *taste* or *smell*—*Infusible*—Spec. grav. 4,84—With *Borax* melts into a white transparent glass—Soluble in *Carbonate of Ammonia*, though not in either of the *caustic fixed Alkalies*—Insoluble in *Water*.

With the *Sulphuric, Nitric, Muriatic*, and with other acids, forms compounds remarkable, like those of *Glucina*, for their astringency and sweetness, to which earth it is considered in many respects as analogous, though in others essentially different.

OF COMBUSTIBLE SUBSTANCES.

The substances usually denominated *Combustible*, are such as are more especially remarkable for exhibiting the phenomenon of *Combustion*, when heated to a certain degree in contact with *Air*.

The changes produced on the *Air* in this process, and the alteration which the *Inflammable Substances* themselves undergo, already particularly treated of.

In their *Form, Consistence, Weight, Volatility, &c.* the substances of this class differ widely

is more divided the Particles of a Soil are
better - Thus Plowing & Harrowing are
good.

Layer when mixed with

The Fertility of Soils influenced by Temperature
at which water the the most is lost

Consequence of the same

Phosphates of Lime has been found in And,
Dens however the same

Ammonia has been found for
many years but the same is not to be seen
lost there is an Elasticity of the
of the soil which can contain a great
quantity of the same

Vegetable matter containing much

Rock being very quickly permeated by
decomposition of hydrogen below in
Purman's Bank. & whenever a hole
contains the roots of Plants as Oak
Rock being very rapid

Carbonate of Magnesia is Purman's
Rock. It is a crystalline rock. This is
indicated by its white & being quickly
large quantities of Carbonic Acid

That earth which is

Vegetable matter & decomposition of Plant
Purman's Bank, which is
of Plants as the Tuffaceous & Purman
There is such a thing as a very good
-rock Pottery is a Purman's Bank
It is of very hard grain & is at
Purman's Bank. Free

on Volcanic formations

found in the *Stratigraphical* for
age

And I am has been found to contain the
following constituents in viz
50 Parts of Fine Sand 5 P. Carbon of same
part of good Plank 15 P. of Silica 14 of Argon
the rest of the Iron

Barren Soil -

00 from Sulphur of Iron numerous
Insects from It from Chalk Limestone
Clay -

Carbon is widely deposited in the globe
Animal & Vegetable Bodies

The Chalk is quite hot the whole is covered
When cooled it makes the fire for use
It can not be found anywhere

Carbon.

It is not found in a solid state but usually
largely in a gaseous form, it exists largely in
Coal Wood &c., etc. dissolved in Air in
form of Carbonic Acid Gas.

Charcoal differs in Purity according to the manner
it is produced.

The Charcoal from Animal Subst. requires a higher
temperature to burn it than that of Vegetable

Common Charcoal is generally produced from
Oak - Chestnut &c. cut up & dried for 2 or 3 months
in place in the ground is prepared & the wood piled
up with alternate layers of brush and holes &c.
then it is fired & the fire is applied to it.

In Scotland they carry on the process of making
Charcoal very extensively.

The Finest Carbon is perhaps procured from
Lamp Black, for very delicate experiments.

Charcoal has a disposition to Absorb Water
from the Atmosphere and increase in Weight.

Charcoal from fine Lignum Vitae Oak &c. making
as a substitute for the finest

from each other. They are comparatively *lighter* than most other bodies.

May be divided into two classes—*Simple* and *Compounded*.

The simple combustibles are, *Hydrogen*, *Carbon*, *Sulphur* and *Phosphorus*.

Hydrogen.

Already treated of under *Aeriform fluids*.

Carbon.

Obtained in its common form *Charcoal*, from vegetable, animal and bituminous substances by exposure to heat in close vessels.

Varies in its form, quantity and purity, according to the nature of the substance from which it is prepared.

The charcoal of common wood, *black*, *light*, *brittle*, *sonorous*, *insipid*, *inodorous*, and of great *durability*.

Capable of sustaining the most intense degrees of *Heat* in close vessels, without alteration; but

upon access of *Air*, burns with a white flame, and yields *Carbonic Acid Gas*, leaving behind a small quantity of *earthy saline Ashes*.

Decomposes the *Sulphuric Acid* and all its compounds, by the assistance of *Heat*.

Decomposes also the *Nitric Acid* without *Heat*, and sometimes with such rapidity as to occasion *Combustion*. Detonates with *Nitre*.

Dissolves, by fusion, in the *fixed Alkalies*; very readily also in *Alkaline Sulphurets*.

Combines by cementation with *Iron* either in a small proportion, *Steel*, or in a larger, *Plumbago*.

Of the properties and uses of *native Plumbago*.

Charcoal remarkable for correcting *Fætor* and depriving many substances of *Colour*, especially when used in its fresh burnt state.

Exists in its present form in the *Diamond*.

Used chiefly as an article of *Fuel* and in certain arts, trades, and manufactures.

Of Sulphur.

Found either uncombined, as deposited by water, or sublimed by subterranean fire, *Native Sulphur*; or in combination with other bodies, more especially with different *Metals*.

For very Delicate Experiments are enclosed
pieces of Box Wood in a Crucible filled
with sand and apply a steady heat for
an hour or more when you will procure
pure Carbon, it should be kept in Dry
stopped Closely, glass Bottles.

The Charcoal of common wood is black, light,
and porous.

get rid of the substance which forms the Charcoal within
and of heat then when made in a crucible
it is porous.

now expose a solution of Silver or Gold to air & heat
the Charcoal coated

Carbon is infusible in any common Heat

Charcoal & Animal Matter when mixed form a very
good lining for Crucibles.

Charcoal is a very durable it is not capable
of Putrefaction: hence the practice of
burying it in the states before they are
ever out of ground.

The Birds don't
eat upon it but it readily decomposes

him. Charcoal is capable of Combining
with iron, it is combined in Black Lead

which being enclosed in Cadaver forms the
Black Lead Pencil. Phosphorus is found in Charcoal

Charcoal detonates with $\frac{2}{7}$

The Best kind is produced in this Country

The Substance of Carbon is Spongy & porous.
it will absorb Carbonic Acid from Water completely

Part of Charcoal in a few hours will absorb
three times its Weight of atmospheric Air

Charcoal very readily will take the coloring
matter of various substances

Charcoal has a great power in correcting taste
of bad Water or Meat, by either throwing it
into the Water in which the Meat is lying
or the Water that has been tainted

This rarely if ever found. Pure. The Diamond
is found to be the most. Pure: it will
withstand the greatest heat.

We are at a loss to know How the Diamond
is formed. They are generally found

Diamond of all known bodies has the
greatest power of refracting the Rays of
light. They are exceedingly hard. The
Diamond is the hardest substance known.

Some Naturalists said that they
were not at all acted on by heat.

When they are colored, it lessens very much
the value, but it is partly got rid of by
applying heat in a close vessel.

Diamond is a non conductor of heat.

The value of them increases very much
with the weight of them.

The largest Diamond is in the
Crown of the King of France.

It is made by heating in close vessels
common Bituminous Coal.

It has been known to grow
in the Earth.

Charcoal forms a very excellent Dentifrice
- a very fine powder.

It is very hard & very
brittle.

Compared with the
Diamond.

Sulphur

Its Specific G. in its native state is 2000 water
being 1000 - when heated 2 G. in 1000 water
is dissolved

The compounds of Sulphur without Oxygen
are called Sulphurets, its compounds with
Oxygen being Sulphates, Sulphur begins
to attract Oxygen at 100° of Fahrenheit than it begins
to melt

The Acid: Nit. acts on Sulphur: so the Chlorine G.
with which it forms a red fuming
Gas.

Sulphur begins to attract Oxygen at a temperature
of 100° of Fahrenheit
at 500° of Fahrenheit it becomes an elastic fluid.

Sulphur is insoluble in Spirits of Wine but in
strongly alkaline

100 Parts of it consist of 15 Carbon less than 0.1%
of Hydrogen & 84.9% of Sulphur

May be obtained artificially by the decomposition of *Sulphuric Acid*.

Colour, *pale yellow*.—Sp. Gr. 2.033.—*Hard, brittle, insipid, insoluble*.—Very easily melted.—Sublimes in close vessels into *light yellow* flowers, formerly called *Flores Sulphuris*.

Burns with a *blue* flame, and by absorbing *Oxygen* from the Air, is converted into *Sulphureous Acid Gas, Sulphureous* or *Sulphuric Acid*, according to the proportion of this principle.—Similar effects are produced on it by deflagration with *Nitre*.

Is little affected by *Acids*.

Unites readily with the *Alkalies*, and also with the *saline Earths*, producing liver-coloured compounds (*Sulphuret of Potash, &c.*) which are soluble in *Water*, and may be decomposed by all the *Acids* (*Sulphur præcipitatum P.L.*)

When intimately mixed with certain proportions of *Potash* and *Nitre*, produces a compound which when gradually exposed to heat, explodes with great violence (*Pulvis fulminans*).

With *Nitre* and *Charcoal*, in mixture, it constitutes *common Gunpowder*.

Combines by fusion with all the metals except *Platina, Gold, and Zinc*.

Is insoluble in *Spirit of Wine*, but unites with oily matter of every kind, and with all the liquid bituminous bodies, into compounds of increased consistence (*Balsams of Sulphur*).

Order of attraction in the moist way, *Lead, Tin, Silver, Quicksilver, Arsenic, Antimony, Iron, Potash, Ammonia, Barytes, Lime, Magnesia, Unctuous Oils, Essential Oils, Æther, Spirit*; in the dry way, *Potash, Soda, Iron, Copper, Tin, Lead, Silver, Cobalt, Nickel, Bismuth, Antimony, Quicksilver, Arsenic*.

Employed principally in *Bleaching*, in the manufacture of *Sulphuric Acid*, and of *Gunpowder*, frequently also in *Medicine*.

Of Phosphorus.

Obtained, by decomposing calcined *Bones* by means of diluted *Sulphuric Acid*, evaporating the supernatant liquor to the consistence of a syrup, mixing it with powdered *Charcoal*, and distilling in the open fire; or by adding *Nitrate* or *Acetite of Lead* to common *Urine*, collecting the precipitate, mixing this with *Charcoal* and distilling as above.

is extremely Volatile, it will decrease the Weight of the Ther:
considerably. it has somewhat of an offensive Smell
as off the hand like Ether when applied -

If you add a little of this fluid to Oxygen Gas
it will expand and by applying Heat
will detonate violently -

Nitrous Acid Gas is produced by applying
Heat to a beautiful flame -

Sulphur when in the composition of gun
powder

Sulphur is found in Sicily in its purest State.

Sulphur has a great effect of restraining the circulation
of these Poisons. When Sulphur is
combined with Alkalies they are termed
Sulphurets.

Common Flamm of Sulphur is commonly known

as a kind of ^{Carbonate of Potash} ~~Carbonate~~ of Potash

Sulphur and gun powder obtain their

name from Sulphur and Potash. If any Acid is

added to it it gives off Sulphuretted Hydro Gas. The Sulphur

is put into a glass

By throwing Sulphuret of Potash into Water &
effect will be

Sac Sulphur is Made by precip^a Sulphuret by
an Acid.

By taking a Solution of Sulphuret of Potash &
by pouring into it Sulphuric or Nitric Acid
and you will form The Sulphur Precip^a

Sulphur is Capable of combining with the Air & also
with Iron & other Metals

It should be kept in Green Bottles

Any Matter is Capable of Disolving and Sulphur
by Boiling it in Olive Oil you form
Balsam of Sulphur. —

Sulphur is widely diff^d thro^ug^h the Earth and
the Class of Veget. called Felted grasses
United with Hydrogen in many Mineral
Waters. . .

a Mixture of 3 Co. Sulph. & 1 Co. Potash. Care
must be taken 3. Ours the which is
used when heat is applied The Cause of
affection is not exactly ascertained.

The Sulphur containing Alumina

to 100 grains of Sulphur in a flower
not with the acid of sulphuric acid
it is deposited under the alumina

effect of Sulphur is not temporary
has an effect of Alumina under
an effect of Sulphur

When the Sulphur is not temporary
it is not the form of Alumina
it is not the form of Sulphur
it is not the form of Sulphur
it is not the form of Sulphur
it is not the form of Sulphur
it is not the form of Sulphur
it is not the form of Sulphur
it is not the form of Sulphur

Sulphur will combine with Phosphorus

Phosphorus

was accidentally discovered in 1777 -

There is a small amount of Phosphorus in
the bones of man & other animals
which is excreted in water. J. G. 1776

Phosphorus is undergoing decomposition when
exposed to light & air - at a temp. of 110° of Fahr
it melts. ^{210 to 220 at 225 it boils} In the steam bath Phosphorus
will be exposed in water to combine with
Oxygen and is found. Capable of uniting
with 3 parts of Oxygen.

Phosphorus is a highly inflammable substance
but the pure form is not so inflammable as the
impure. Phosphorus is melted by heat & is superior to
all other substances in which Phosphorus is found.
It should be as much as possible of the pure
form. It should not keep a large quantity in water as
in cold weather the water may freeze & the
phosphorus will be exposed to the air & will
be oxidized & become a phosphine.

Phosphorus may be prepared in many ways
by the action of acids.

When an acid solution of Phosphorus is exposed to the air
it becomes luminous.

At the temp of 100° it burns & at a hundred & twenty it
burns very quickly.

Purified by cautious re-distillation, or straining it, when melted, through leather.

Colour *pearly-white*. *Semi-transparent*. *Waxy*.
Insoluble in Water. *Very fusible*.

When exposed to air, at a low temperature, emits a white fume, and is luminous in the dark; if heated, burns with great rapidity; and in both cases acquires acid properties,
Phosphoric Acid.

Decomposes the *Nitric Acid*, occasioning combustion by the sudden separation of its *Oxygen*.

By treatment with the *fixed Alkalies* or *Lime*, yields a permanently elastic fluid which explodes on admission of *Air* (*Phosphuretted Hydrogen Gas*.)

Dissolves in *Essential* and *Unctuous Oils*, *Spirit of Wine* and *Æthers*.

Unites by fusion with *Sulphur*.

Unites also with several of the *Metals*, and decomposes most of their *Oxyds*.

Used in the preparation of *liquid Phosphorus*, *Phosphoric Matches*, and portable *Phosphoric Bottles*. Attempts have also been made to employ it medicinally in small doses.

Compound Combustibles.

Found abundantly in the vegetable, animal, and mineral departments; some prepared artificially.

Consist principally of *Hydrogen* and *Carbon* variously combined.

May be divided into five kinds, *Alcohol* and *Æther*, *Fixed Oils*, *Volatile* or *Essential Oils*, *Resin* and *Bitumen*.

Of Alcohol.

Obtained from such organised substances, or their products, as have undergone the *Vinous Fermentation*, of which those containing *saccharine Matter* are alone susceptible.

This process materially influenced by *Rest*, *Dilution*, *Temperature*, and *Exposure to Air*.—Divisible into different stages, the *vinous* productive of *Alcohol*, the *acetous* affording *Vinegar*, and the *Putrid* generating *Ammonia*.

These changes promoted or retarded by various means, *Ferments*.

The lastest I saw a quantity of iron which is
imported to the Government by a Lyceum
must be the same as that which I have
before given me a picture of Phosphorus.

... than look in another which
which ought to be look I must have I
... ought to be made at the
mouth, the mouth of the mouth should
put into water

ought to be of about the 3 part of the
elf I don't know it on the other
is it hard for a few days then get
by inspection you

the glass of Phosphorus is being made it is
little channels & put into a water

the wine must be quite brown & must be
by the head & by the tail


Alcohol

... is always attended with more or less
... all substances containing sugar are
... into various degrees. These are
... which contain a
... matter

If sugar is fermented with about 1/2 of alcohol
... by abstracting a portion of the
... The lower the temperature when
... is going on the more spirit will
... During the process
... is evolved.

... is usually contained
... by the acid
... is proved by the

Alcohol is very inflammable and burns
up residues, if you put a little Gun
powder into a small quantity of Alcohol
and set it on fire, the gunpowder after
the Alcohol has burned off the good, will
explode.

The diagram illustrates the process of distillation or separation. It shows two circular vessels, likely retorts or flasks, connected by a horizontal tube. A vertical line passes through the center of each circle, possibly representing a central axis or a point of measurement. The text above the diagram mentions 'distillation' and 'separation'.

Alcohol readily mixes with Water in any
proportion, with a diminution of volume
and an increase of heat. Showing that the
two fluids have a strong attraction for each
other. By pouring Alcohol on a solution of
Sulphate of Potash, the salt will be precipitated.

Muriate of Lead. Silver Oxide insoluble in
Alcohol.

Sulphate of Lime or Strontian, is quickly
precipitated by alcohol. When dissolved in water
Alcohol is decomposed.

It does not act on Sulphur except in a state
of vapour. Both of them very little affected
by Alcohol.

Fixed Oils as Olive Oil will not combine with
Alcohol, but it will, when its constituents are
Alkali. The flame of Alcohol is very much
increased by depositing Colours with the Alcohol. But
Alcohol is also a powerful Antiseptic
and used for the purpose of keeping Anatomical

Repeated distillation and digestion on dried
Muriate of Lime or on Potash, necessary to
bring Alcohol to its utmost degree of purity.

When pure colourless and transparent.

Taste hot and pungent. Sp. Gr. 0.815.

Miscible with Water in all proportions.

Burns with a bluish flame, producing in com-
bustion Carbonic Acid Gas and Water.

Undergoes singular changes in its properties
by treatment with the different Acids; hence
the preparation of Æthers, Oleum Vini, &c.

Dissolves the Alkalies, and many of the
neutral Salts, particularly such as are deliques-
cent. Also dissolves Soap, and acts readily
as a solvent on Essential Oils and Resins, on
Balsams and on Camphor; hence the prepa-
ration of various Spirituous Liquors, Tinctures,
Varnishes, &c.

Considered by Lavoisier and others, as a com-
pound of Hydrogen, Carbon, and Oxygen, pro-
duced by the decomposition of the sugar, in the
act of Fermentation.

Order of Attraction, Water, Æther, Essential
Oils, Ammonia, Potash, Alkaline Sulphurets,
Sulphur.

In general use for various technical as well as
dietetical and medical purposes.

The Hydrometer is the instrument used to
ascertain the specific gravity of alcohol
and the test is weight 700 grains the
of weight in grams is light in covered tank

Of Fixed Oils.

Obtained both from *Vegetables* and *Animals*.

From the *former* sometimes by boiling, but mostly by pressure, from certain *Fruits*, *Kernels*, and *Seeds*, hence called *Expressed Oils*; called also *Unctuous Oils*.

From the adipose membrane of the *latter*, of different colours and consistence according to the part, the age, and the species of animal from which it is taken, *Lard*, *Suet*, *Fish Oil*.

The difference between *Animal* and *Vegetable Oils* not exactly ascertained. The *Animal* in distillation furnish a peculiar acid, *Sebacic Acid*.

Both vary in the temperature at which they become *solid*; are insoluble both in *Water* and *Alcohol*; become *rancid* by keeping; yield, by distillation in close vessels, an *acid Phlegm*, a lighter and a denser *Oil*, (*Empyreumatic Oil*), a large quantity of *Hydrogen Gas* mixed with *Carbonic Acid Gas*, and leave behind a small proportion of *Charcoal*.

Afford *Water* and *Carbonic Acid Gas* by inflammation in contact with *Air*.

By mixture with the stronger *Acids*, produce,

Sulphure Ether is made by taking
equal parts of Alcohol & Sulphuric Acid
then must be mixed cautiously & then
not applied very gently.

As it is obtained in $\frac{1}{2}$ first instance
contains Sulphurous Acid, when you

then contains a greater proportion of
Sulphur, and a smaller portion
Carbon. Ether will not boil till
temperature of 200 of Fahrenheit: Thermom.
Sp Gravity of ether

The latter stage of $\frac{1}{4}$ Proof we have
of wine,

may obtain the Al Vin; by taking in
first instance a larger portion of Sulphur
in $\frac{1}{4}$ Proof - when you will have
a little Ether but a large quantity
Al: Vin

Small portion of Ether may be formed without
aid of heat by a mixture of

Aether will not readily unite with Water
which may be observed by lighting it
on the surface of Water.

This one of the Lightest & fluest we have
It boils at 60 in the air & 20 below Zero
in a Vacuum. You & Aether evaporates
very quickly at 60 degrees below Zero in the
Air. ~~Chlorine~~
it is quickly decomposed by Chlorine, and

Oxygenized Muriatic Acid.

When the Aether is good, by putting a small
of it, into Chlorine it will slightly detonate and
The inside of the vessel is covered with Carbon
This experiment should be made with Caution
DANGER is sometimes the result of it but not ordinary
but this ought to be done carefully.

Aether has $\frac{1}{4}$ property of dissolving a portion
of Gaseous

Aether is also obtained by mixing Alcohol
with Nitric Acid, but this should be done
with Caution.

It is extremely Volatile so that if exposed in Water
Whether you will observe a number of air
bubbles.

Nitric Aether also has the power of dissolving
Bodies.

Other: Nitroas: Made by 4 Parts of Alcohol and
 part of strong Nitric Acid by weight
 mix them continuously & draw off 3 Parts & pour
 what remains a good deal of brown material
 spent other still more by exposure to air get gentle Acid
 By mixing Alcohol 2 Parts by measure
 and one part of Nitric Acid, this will
 have the action of the Nitric Acid
 & Alcohol.



Nitrous Ether
 beverage

Munatic Other is not
 found by employing the Smelling liquor of Am
 Alcohol Take walrus off put Alcohol into them
 in with common Salt & remainder. Sulf Acid

a very agreeable Other is
 Prepared by equal parts of Acetic Acid
 & Alcohol - distilled with gentle heat
 pendant having a good deal of Acetic Acid though
 it is up with Carbonate of Soda

As general properties, The Acetic
 the, resembles very much the other
 but is employed much in
 medicine.

Fixed Oils

Some of them are dissolved under not long contact at an
common temperature of the atmosphere

Generally less than that of water

Others are perfectly soluble in Alcohol from free
water which may be prevented by the addition of
little camphor

Some are unchangeable & others when exposed to the air
they become thick & soapy these are called fatty Acids
because abstracting hydrogen from them. In others the
the property of hardening them from contact with
Acids as the oils from various seeds. The fatty Acids
by long exposure become drying by losing the
hydrogen volatile in fatty Acids

Sulphur readily separated in fatty Acids
And Sulphur has considerable action on fixed
If you take fatty Acids slowly burning & drop a
concentrated sulphuric Acid on them you
will observe a white or light brown solid mass
formed

None of the volatile Salts decompose
These Oils when exposed to the atmosphere when rubbed
with finger or with paper, Fixed Acids are
formed or being transformed from the air

in some instances, *saponaceous Compounds*; in others occasion *Combustion*.

Unite more perfectly with the *Alkalies*, more especially with *Potash* and *Soda*; with the latter, form common *Soap*.

Unite also into saponaceous compounds with *Barytes*, *Lime*, and *Magnesia*.

Have no action on any of the *Metals* except *Copper* and *Iron*, but assisted by heat dissolve most of the *metallic Oxyds*, and when separated again from these, are found to be soluble in *Alcōhol*.

Combine by the assistance of heat, with *Sulphur* and with most of the *Bituminous Bodies*.

Considered as differing principally from the *Volatile* or *Essential Oils*, in containing different portions of *Mucilage*.

Employed in *Painting*, *Varnishing*, *Soap-making*, in *Mechanics*, for *Fuel*, in *Diet*, *Medicine*, &c.

Of Sperma-ceti.

Obtained from the brain of a particular species of whale, (*Cetus dentatus* Lin.) thence called the *Sperma-ceti* whale; obtained also in small quantity from the oil of the same fish.

After refinement, *white, semi-transparent, crystalline, friable, insipid, inodorous.*

Differs chiefly from the fixed oils in being little altered by distillation, and not easily acted upon by *Acids* or *Alkalies*.

Supposed to have the same relation to *Unctuous Oils* that *Camphor* has to the *Essential*.

Used chiefly for making *Sperma-ceti* candles, and in the composition of *Ointments, Plaisters, &c.*

Of Bees-Wax.

Deposited by the *Bee* in the construction of the *Honey-comb*. Colour, *yellow*.

Very analagous to *Fixed Oils* in all its essential properties.

Becomes white on exposure to *Air*, or treatment with the *Oxygenized Muriatic Acid*.

Forms the basis of several *Cerates, Ointments, and Plaisters*.

Employed also in several of the *Arts*, but principally in making *Wax candles*.

Of Volatile or Essential Oils.

Obtained from most fragrant vegetables, by expression, or distillation with *Water*.

Alkaline solution the stronger you throw
the 24 Oz of Salt 12 Oz of Lime in
solution of water heat them & evaporate
until you make a bottle that will hold
a pound of water since hold on 24 Oz of
the same

boiling water soap in them cannot be alkaline what
of for Heat? Soap is made by the solution
of Alkali and oil.

Soap water is not made in Cold Soap

as being too readily decomposed by an Acid
than common hard Soap there is in Water of
lime in the preparation of the

for Soluble in Alcohol. 12 Oz of Soap is often
employed for a Chemical Test as for Soluble
in water is said to be harder than Saponified.
the same being exactly salt to which many
are added by in the Cold Soap.

in kinds of Soap as Rotten which is Down
many of some of the other Oxydation on the
Oxydation of Soap

any preparation that just Oil contains

expensive and also from being generally got
from India. Sometimes found in the
leaf of plants. Known as mother of pearl
very little.

Ground Air may be found. It is found in
the bottom of the sea. The best known
is the mother of pearl. It is found in the
leaf of plants. It is found in the
leaf of plants. It is found in the
leaf of plants.

Ground Air is obtained by Professor. It is found
in the bottom of the sea. It is found in the
bottom of the sea. It is found in the
bottom of the sea.

The difference of the air is dependent on the
method of getting it. The ground air is
found by using the mother of pearl. It is found
by using the mother of pearl.

air from Poppy seed which are a source of
dry air. It is found in the bottom of the sea.
It is found in the bottom of the sea.

It is found in the bottom of the sea. It is found
in the bottom of the sea. It is found in the
bottom of the sea. It is found in the
bottom of the sea.

It is found in the bottom of the sea. It is found
in the bottom of the sea. It is found in the
bottom of the sea. It is found in the
bottom of the sea.

The Balsams some of them have a pleasant smell
and some are very good for the skin
The Balsams some of them have a pleasant smell
and some are very good for the skin
The Balsams some of them have a pleasant smell
and some are very good for the skin

Copal is a species of resin but this not soluble in
Alcohol, but like the other Resins it is soluble in
alkalies and Nitric Acid. The best Copal comes
from as it is not soluble in Water
and very little to other Substances, it forms
a good Varnish, for Preparations for

3lb of Powdered Copal 3oz of Al: Turpentine
apply heat until the Al: Turpentine boils
then add gradually the Powdered Copal until
you find the last portion is hardly acted upon
at all the strain it out and then let it settle
the last Al: Turpentine
Powdered 3lb 3oz of Al: Turpentine 3lb
Powdered 3lb 3oz of Al: Turpentine 3lb
Powdered 3lb 3oz of Al: Turpentine 3lb

Taste *pungent*. Odour, colour, and consistence, *various*. Most of them lighter, some heavier than *Water*.

Thicken and become less odourous by absorbing *Oxygen* when exposed to *Air*; hence the supposed formation of *Balsams* and *Resins*, which differ chiefly in consistence and the occasional presence of *Benzoic Acid*.

Also affected by exposure to *Light*.

Sparingly soluble in *Water*; but readily so in *Alcohol*.

Volatile in close vessels, at or under the temperature of 212 *Farenh*. By repeated distillation lose their characteristic properties, and are brought nearly to resemble each other.

Highly inflammable when heated in contact with *Air*.

Decompose the stronger *Acids*; in some instances with such rapidity as to occasion actual *Combustion*.

May be united with the *Alkalies* and *Lime* so as to form *Soaps*.

Combine intimately with *Sulphur*.

Combine also with *Phosphorus*, *Unctuous Oils*, and *Camphor*.

Purity ascertained by solution in *Spirit*, or exposure to *Heat*.

Chiefly employed in *Medicine*.

Of Camphor.

Thought to be a *Volatile Essential Oil*, containing a large proportion of *Carbon*.

Exists in many of the fragrant plants, as *Lavender*, *Rosemary*, *Marjoram*, &c. but principally procured by distillation with water, from a particular species of laurel, (*Laurus Camphora* Lin.)

Requires certain additions in its subsequent refinement by sublimation.

White. *Transparent*. *Friable*. Taste pungent and *bitterish*. Specifically lighter than *Water*.

Evaporates completely if kept exposed to the *Air*.

Burns with a white flame, and is entirely consumed.

Sparingly soluble in *Water*; but readily so in *Spirit of Wine* and *Æthers*, in *Unctuous* and *Essential Oils*.

Dissolves both in the *Sulphuric* and *Nitric Acid*, without decomposition; by repeated dis-

Opal in many of its properties resembles
amber. Amber

Camphor

Camphor

Camphor is found interspersed in the veins of the
phosphoric acid; than essential oils

Boiling point 302 Deg. of F. It
if heated suddenly
Camphor is highly infl. Carbonic Acid & peroxide
Camphoric Acid

Lighter than water, it is easily burned
in air.

in warm weather it evaporates completely
It is very inflammable, and leaves no residue
has been in years of Carbonic Acid &
phosphoric Acid.

By repeated Distillation with Nitric Acid Camphor
is converted into a peroxide of Nitrogen

Alcohol Ether & Oil will dissolve it.
In Alcohol, precipitated by water, the
water having a greater attraction for the Spirit
than $\frac{1}{4}$ Camphor does.

There is a kind of Camphor in Japan which
under goes Camphor

Camphor has been found in many Plants
as Laurander Rosemary &c

Water with only $\frac{1}{10000}$ ^{the} Part of
abundantly soluble by alcohol

By the intervention of Gum or Sugar Camphor
with water can form an emulsion

If a small piece of Camphor is placed on Plain Water
It will begin to move round the surface very gently
but if any portion of essential oil is added it
motion will be rapidly stopped this has
been satisfactorily explained.

By passing through a retort open to the air the volatile
oil of asphaltum (asphaltum) is formed.

Bituminous Bodies

They are either liquid or solid & found
in many degrees of Purity. Kerosene is
one of the most pure. It is found in
Naphtha... is one of the most pure. It is found
at Baku in the Borders of Caspian
is also in Italy & other places. Other

It is composed of Carbon & Hydrogen, simple
proportion to the Air will carry off a large
portion of Hydrogen.

Petroleum or Tar

As the Carbon is increased it will more
readily combine with Alcohol

Mineral Pitch or Maltha

Resembles Common Pitch—

distilled harder it is called Asphaltum
which will Melt & is very inflammable

The Main bulk of It is Asphaltum
It may be burned into various Shapes
as is specified in Making ornaments

Canal Coal, is full black

Produced in Lancashire

100 Parts contain 72^{pts} of Carbon 22 of Maltha
(but no Asphaltum and earthy Matter)

When Distilled it affords an oil like Turpentine
and what is left will be Coke—
which contains more Carbonaceous Matter & is
better than of Canal..

100 Parts of Whitkham, ^{Parts of} 57 Carbon 41 of Maltha & Asphaltum
and nearly 2 Parts of earthy residue—

tillation with the latter, is converted into a peculiar *Acid*. Artificial substance resembling *Camphor*.

Chiefly employed in *Medicine*.

Of Bituminous Bodies.

Found either liquid and devoid of colour, *Naphtha*; or more or less liquid and of a dark colour, *Petroleum*, *Barbadoes Tar*; of a harder consistence, *Maltha*; or perfectly solid, black, and of a compact or slaty texture, *Asphaltum*, *Jet*, *Coal*.

The more liquid *Bitumens* by continued exposure to *Air*, convertible into the more consistent, and all yielding similar products in distillation. *Coal* considered as a variety of *Bitumen*.—Origin and formation of *Coal*.—Artificial formation of *Bitumen* from vegetable Matter, by heat and pressure. In the different kinds of *Coal* there exists more or less earthy admixture, and often *Iron Pyrites*.

The *bituminous* bodies immiscible with *Water* and insoluble in *Spirit of Wine*.

Of Amber.

Nearly allied to the foregoing.

Found for the most part in irregular masses more or less transparent, and of a brownish or yellowish colour.—Specific gravity from 1.055 to 1.000. Emits a peculiar odour on friction and becomes *electric*. Melts at 550 *Farenh.* Burns with a whitish flame. In distillation yields *Water*, *Empyreumatic Oil*, (*Oleum Succini P. L.*) and a concrete *Acid*.

Insoluble in *Water* and nearly so in *Spirit of Wine*, also in all the acids, the *Sulphuric Acid* excepted, in the solutions of the *Alkalies* and in *essential* and *expressed Oils*; but the *Balsams* dissolve it readily.

Of the methods usually employed for rendering *Amber* transparent.

Is probably of *vegetable* origin.

*Antique Amber is prepared by taking
a quantity of Amber and placing it in
a vessel of water and boiling it for
several hours.*

Amber

is an imp. substance it is a brittle transparent
solid substance

Specific gravity little more than that of
water. when melted in various substances

Water does not act on it, but alcohol will
solve for part of it. Alkalies will
solve it also. Nitric Acid... it must be
-treated considerably, like Copal before
it can be used for a varnish. But
it ought to be ~~carefully~~
of Roasted Amber 3 24 of Turbenth.

The Acid. afforded by the Distillation of
amber already spoken of. is called

is Saccharum affords also Ol. Succini
prepared by Distilling this it becomes

amber abundantly found in Persia
India & eastern shores also found

It is probable that Amber is formed from
volatile substances which have been acted
- by Sulphur

the air & acid used to be used in medicine

^{signified} This is a valuable essential combined with
a large portion of Carbon Hydrogen & Oxygen

Coal

Coals are found in the earth seldom
in strata but in seams.

The thickness is mostly uniform. The direction
of them generally east & West.

The seams don't run parallel with the
Horizon. The quantity of ^{air}
the deepest seams of Coal: That means by
the surface is commonly the worst.
but this is not always the case.

^{called wood coal}
a species of Coal, is found in Devonshire
more lately; it lies in strata, but
the strata are not parallel to each other
direction from east to west, there is evident
traces of wood in it.

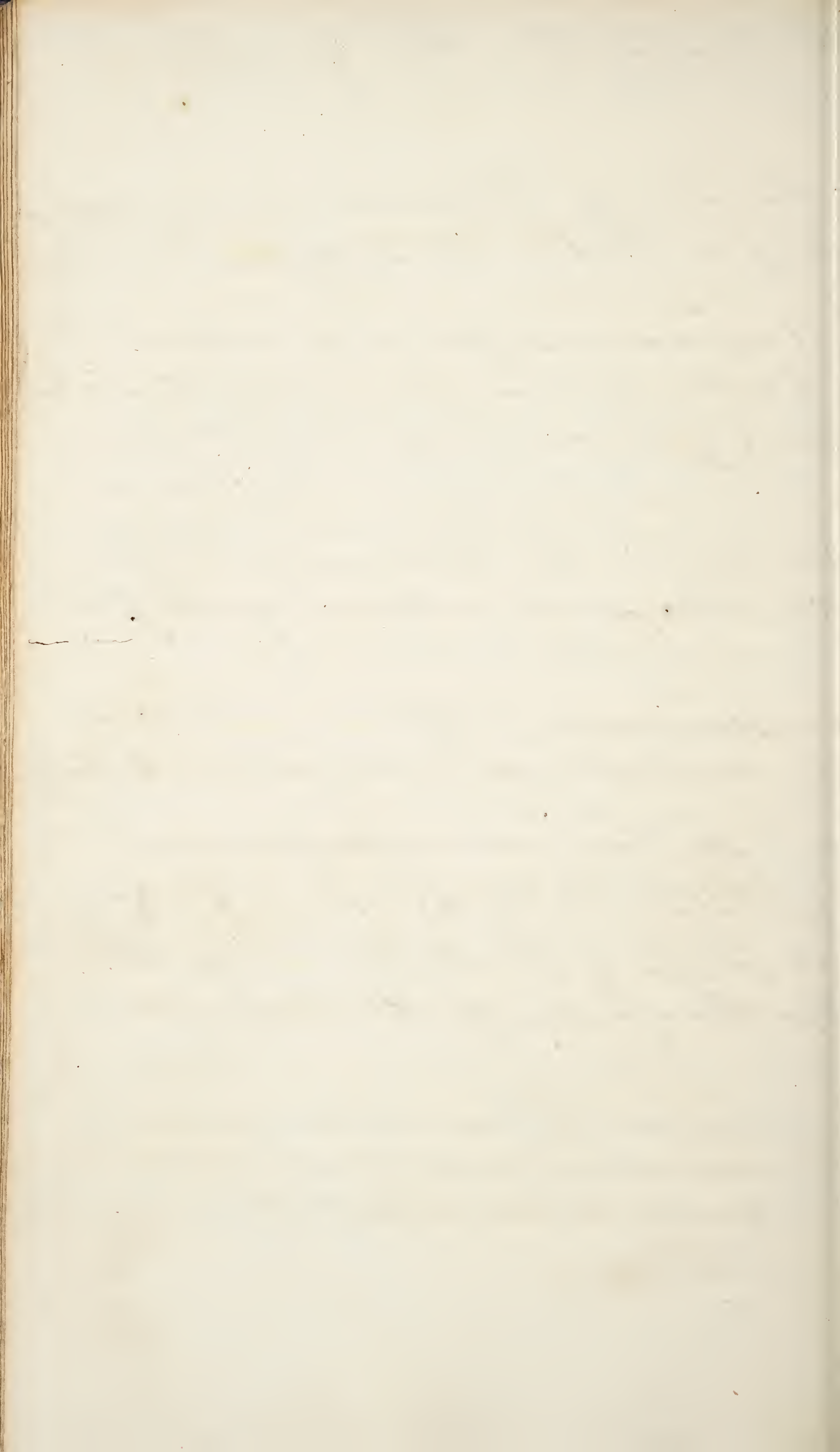
When Carbon has been freed from $\frac{2}{3}$

The character of Petroleum is very apparent
in its nature & heat, the best found in Baffin

Amblygon was formerly supposed to be
a bitumen but now known to be of Animal Origin
its parts of it consists of 5 $\frac{1}{2}$ of Adiposin
, 11 of Benzene Acid 5 of Charcoal
it has been in the stomach of the person
who, considered a disease of this fish
it is inflammable & softens in a moderate
heat, only Nitric acid will act on it strongly
and by some as a perfume.

of Amblygon, 3 of Sugar, a few gr of
oil of Spirit Ben. rectif. This
makes the essence of Amblygon

Essence of Amblygon
Amblygon Acid & Ether also depalms it
in Diluted it becomes fragrant



OF METALS.

Found generally in the clefts or fissures of stony or other strata, forming what are called *Metallic Veins*; or in indeterminate *Masses*; or in *Beds*; or *disseminated* through other substances.

In these instances they are either *native*; or *alloyed* with each other; or in different states of *Oxydation*; or *mineralized* with certain combustible bodies, particularly *Sulphur*; or lastly, combined with one or other of the *Acids*.

Different processes therefore commonly necessary for obtaining them in their separate and proper forms; as, *Pounding*, *Washing*, *Roasting*, *Amalgamation*, *Reduction*, and *Refinement*.

When pure, easily distinguished from other bodies by their united properties of *Weight*, *Opacity* and *Splendour*, as well as by their power of conducting the *Electric Fluid*. Great specific *Weight*, however, is not a character common to all *Metals*.

Some remarkable for their *Ductility*, or *Malleability*, or both; others for their comparative *Brittleness*: hence the common though *inaccurate* distinction into *Metals* and *Semi-metals*.

Differ also from each other in their comparative *Hardness* and *Softness*: when hardened by mechanical extension, may be made soft again by exposure to red heat—*Annealing*.

All *fusible* in close vessels if heated to their respectively necessary degrees; are convex when in fusion, and in cooling disposed to *crystallize*.

Some, particularly *Iron* and *Platina*, grow soft before they melt, and hence are capable of being united by the operation of *Welding*.

Some, as *Platina*, *Gold* and *Silver*, remain *fixed* during fusion; others, as *Quicksilver*, *Arsenic*, *Zinc* and *Antimony* are *volatile* or convertible into a state of *Vapour*.

Variously affected on exposure to *Air*, by which in some instances they are merely *tarnished*, in others deprived of their metallic properties more completely, being converted into a state of *Oxyd*: hence the utility of *Tinning*, *Silvering*, *Gilding*, &c.

All, except *Platina*, *Gold* and *Silver*, still more readily *oxydated* by the united action of *Air* and *Heat*; hence their division into *Noble* and *Base* metals. The circumstances in which these differ from each other in their *Oxydation* are; the comparative facility with which they become *oxydated*, the temperature required for that pur-

to know whether an Ore contains any Metal
by means of a piece of Charcoal, and
putting a little of $\frac{1}{2}$ Ore into a hole in the
charcoal, and throwing a little black flux
and apply heat & you will distinctly
see Metallic Globules.

The Ductility & Malleability differ very
much: the most remarkable Metals for
this property are Lead Iron and
usually Gold & Platinum likewise
are all capable of being drawn out
to a Wire, and beaten into an extreme
thin Sheet. The thickness of Sheet Gold
according to experiment is not found
to exceed one 28000th part of an Inch
thin and Duckslime when cooled to $\frac{1}{2}$
refracting temperatures, will be found
ductile & Malleable.

Antimony Bismuth &c are not found
possessing these properties remarkably.

They differ also in Point of Hardness
as we know even when converted into Steel
in $\frac{1}{4}$ Case of $\frac{1}{4}$ Instrument called $\frac{1}{4}$ file
our Case are alloyed. Thus $\frac{1}{4}$ Silver
Gold can mix with an inferior Metal

exposure to heat. Iron is able to
extend into a sheet many Metals. even
crack if heat was not applied.
This is called by Process of Annealing.
It is often of importance in construction
of various Machines to know, $\frac{1}{4}$ degree
of hardness of a Metal. Dr Babington
has an instrument to effect this
given him by Mr Phipps.

Metallic bodies generally are capable of fusion
but they differ in $\frac{1}{4}$ respect to $\frac{1}{4}$ degree of
temperature, namely when in fusion
and are casting were disposed to
Crystallize. Lead runs fuses much
more easily than Silver but Silver
is more easily fused than Iron
Platinum requires an intense heat to
fuse it.

In cases where solder is applied it might
be that it should be more fusible than the
substance to be soldered.

can't by fused chamber. The
with ductility.

you wish to separate any two
metals, as lead with copper, you
must expose it to a degree of heat

form of metals as Platinum & Iron
now soft before it becomes fused
this is very useful as it renders
them capable of being united by
application of heat. This is done contin-
ually by Blacksmiths. In the
Blacksmith has heated his Iron
to a red which makes it
large. Thus the solder can be
which can unite the metals by de-
to a state of glass.

can't take oxide of Iron, and mix it
with a flux without excluding it from
them, otherwise you will have a
It can be done in a mortar (or close vessel)
Metallic Bodies are affected by exposure
to air Gold Platinum & Silver are not
tied on. but are fused copper Iron
will become affected. Some of
the of Tinning, Silvering and Gilding
which have been are not affected.

We can tell an Iron, Copper, & Silver
When Lead is Melted its surface exposed
to Atmosphere its metal form an Oxide.
Iron may be melted very readily by means
of Oxygen Gas. Copper Burns in Oxygen Gas
with a green flame

The Most effectual modes of Oxidating Metals
is by means of an Acid. Combined with
Water, thus Concentrated Sulphuric Acid
will scarcely act on it, but very strong
when diluted, the reason of this is owing, it
said to be a predisposing affinity, of $\frac{1}{2}$ water
The Acid is $\frac{1}{2}$ Solvent of $\frac{1}{2}$ Oxides not of Metals
When a Metal is brought to a State of Oxide
its capable of forming with $\frac{1}{2}$ Acids, Neutral
Salts

No Metal can be dissolved in Acid excepting when in a State
Oxide -

The ^{Metal} ~~Oxide~~ ^{sometimes} becomes insoluble in Acid by Contact
too much Oxygen - this is the case with Peroxide
Manganic and Molybd. & Chromium are capable
combined with enough Oxygen to bring them to the State
all Metalline Oxides may be may again ^{be made}
liquids, or ⁱⁿ its metalline State.

It may also may be brought into its metalline
State by Chemical affinity. Thus $\frac{1}{2}$ Stront
of Bismuth decomposed by Water its
having a stronger affinity for Acid than
the Bismuth

pose, the quantity of *Oxygen* which they respectively imbibe, the force with which they afterwards retain it, and the phenomena which they present during *Oxydation*.

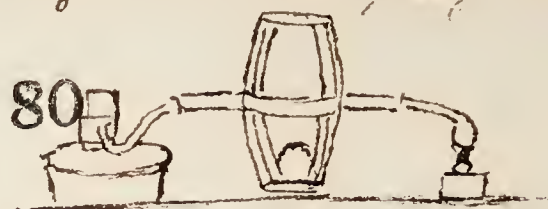
Of *perfect* and *imperfect* metallic oxyds.

Metallic Oxyds, however produced, vary in their properties, not only from each other, but also from their respective metals. Those of the volatile metals much more fixed than the metals themselves:—Some capable of being decomposed or deprived of their *Oxygen* by heat alone, as those of *Platina*, *Gold*, *Silver*, and *Quicksilver*—Such as are not decomposed by the simple application of heat are *vitriifiable*; hence the necessity of *Fluxes*.

Remarkable for the *Colours* and sometimes for the *Opacity* which they communicate to *Glass*: hence the preparation of *Pastes* in imitation of precious stones, and also the preparation of *Enamels*.

Metals also capable of being oxydated by the decomposition of water—Few of them at a common temperature have any considerable action upon *Water*, in general therefore void of *Taste*; some however when aided by a red heat readily decompose it, and are thereby converted into

*Oxydes of Iron & Lead soluble in Soda, Oxydes of Copper
& Cobalt soluble in Ammonia.
Water may be decomposed by being made to pass thro' red iron
as in this Apparatus*



Oxyds: hence the explanation of their effects on various other fluids.

Metals also capable of being oxydated by acids—Previous *Oxydation* necessary to their combination with acids (*Metallic Salts.*) Hence, in their solution, the partial decomposition of the *Acid* itself, or of the *Water* by which it is diluted; and the consequent evolution of *Sulphureous*, *Nitrous* or *Hydrogen* gas.

In some instances they are furnished with so large a proportion of *Oxygen* as to become *insoluble*, in others acidified. (*Acid of Arsenic, of Molybdena, &c.*)

Metallic Salts for the most part possessed of *Colour* and more or less *Corrosive*.

The *Acidifiable Metals*, and those, the *Oxyds* of which have a disposition to combine with *Earths* and *Alkalies*, may be readily oxydated by these through the medium of *Water*.

Many *Metallic Oxyds* soluble in solutions of the *Alkalies*, forming with them permanent compounds. The partial decomposition of the *Oxyd* by the *Alkali* sometimes necessary to this.

Neutral Salts, acted on but slowly by *Metals* unless aided by *heat*, which, by assisting to decompose their acids, oxydates the metal and renders it combinable with the *Alkali*.

Metallie Solution can be decomposed
by Alkalies & the Oxyde precipitated. A solution
of Bismuth decomposed by water alone
leaves a little ammonium on the degree (Oxyde of
Hydrogen) and $\frac{1}{2}$ degree is also become decom-
posed. A Slip of Copper immersed in
a solution of Sol Silver will become covered with
a coat of Quicksilver, If a Slip of Iron be
put into Nitrate of Copper its wire becomes
covered with Copper - It can be removed by
Acid? - there is a proportionate quantity of iron
dissolved in $\frac{1}{2}$ and also throughout there is an exchange
in many metals - coated by Nitrate of Quicksilver when put
in is capable of combining with simple Infl. Subst.
Hydrogen Gas will dissolve of portion of Iron

Steel is a Carburet of Iron.
Metals except Gold & Silver will combine with Sulphur
Uncombined Hydrogen Gas have a considerable
action on Metals, it will tarnish Silver
instantaneously, and to this Gas, is to be ascribed
the ready tarnishing of Silver in Shops
Metals will combine with each other forming
Alloys, which is always harder, ^{by Fusion} than either of
the separate Metals, Soldering depends
on this property of forming Alloys -
is made of Parts, 5 Pts of Lead 3 Parts Tin, these
two together are capable of dissolving in
boiling Water, if Quicksilver be added its more
soluble

The Caustic effect of Oxygen seems to depend on the
supply of free Oxygen to the animal fibres.

The Classification of Hospitals is of very little consequence
The Best is to be divided into 2 Classes those who
are & are not suffering from disease by accident

2 Those which can be deprived of their Oxygen, by carbon or heat alone. The last might be subdivided into the Malleable Metals & Brittle.

Gallium Lind. Mistle Fern Palladium Mercury
 Iron & Chinese arsenic Boron ant^y Mangan
 Tellurium Tungsten Molybdenum Cobaltum or
 Uranium — &

Talks Wallaston he thinks that oxygen enters
into in certain proportion,

Further than last few months experiments have
made to show that $\frac{1}{2}$ same law prevails in
vegetable Kingdom as in the Ales &c

The, Calcium Hydroxide and a considerable amount

The 5 last have now been obtained in their pure state.

The 2nd ~~and~~ they ~~have~~ are very numerous

Metals are employed as the coloring
matter in making Pastes. — The Oxide of
Zinc

Oxides of Metals are poisonous which are
not in their regular state — therefore
the Metals that are poisonous are those
that are affected by the Oxygen

union of Metals with the Oxygenated Acid

There is every reason to believe in our present
state of knowledge that metals are simple bodies

is a question whether oxygen will combine
with metals in determinate proportions only

Platina

In Spanish means Silver, called Platinum by
the Europeans. It is not capable of Oxidation tho
clap'd with Gold by Salts.

Platina has for years been introduced into the Arts

Platina was for a long time supposed to be peculiar
to South America.

The Natural History of it is imperfectly known.

Always found accompanying Gold.

The grains in which it is found are sometimes
coloured bluish.

They are partly attracted by Magnet. This is owing to
portion of iron being mixed with them.

Platina has been found in large pieces, and
in some large masses. It was found in 1763
at Mexico.

Platina in its natural form is superior to
all others. It is not so easily corroded as
other Metals. It is not very much affected
by acids.

Metallic Oxyds already formed sometimes combine with *Neutral Salts*, and produce with them triple compounds.

Iron the only metal capable of combining with *Charcoal*; but this, and almost all the others, or their *Oxyds*, unite readily with *Sulphur*, their common mineralizer: many also combine with *Phosphorus*, and form fusible compounds.

With very few exceptions they unite with each other by fusion in all proportions.

Metallic bases of the earths and alkalies. General history of that discovery.

Classification of metals.—The metals at present known are the following; *Platina*, *Gold*, *Silver*, *Quicksilver*, *Lead*, *Copper*, *Iron*, *Tin*, *Bismuth*, *Nickel*, *Arsenic*, *Cobalt*, *Zinc*, *Antimony*, *Manganese*, *Wolfram*, *Uranium*, *Molybdena*, *Titanium*, *Tellurium*, *Chrome* and *Columbium*.

Of Platina.

Brought from *South America*, in small grains of a dull silver colour, and commonly mixed with

Quicksilver, Ferruginous Sand, particles of Native Gold, and no less than four new discovered Metallic Bodies. Found also sometimes, though rarely, in lumps—Bed and matrix unknown.

Manner of purifying, by fusion with *Phosphorus*; or amalgamation with *Quicksilver*, and solution in *Oxy-muriatic Acid*—Wrought in the large way by repeatedly melting with *Arsenic* and *Potash*, and subsequent roasting and hammering.

When pure, of a *Silver white* colour, inclining to *Iron-grey*.

Malleability and *Ductility* intermediate between those of *Gold* and *Silver*—*Hardness* greater than that of either—*Tenacity* next to *Copper*—*Specific Gravity* in its native granular state 17,7; in its malleable state 21,3, but varying a little according to the degree of compression.

Extremely difficult of *Fusion*; but when urged by an intense heat, becomes capable of being welded, though imperfectly.

Less disposed to *Tarnish* than either *Gold* or *Silver* on exposure to air—and not *Oxydable* by it even under the strongest heat; but may be reduced to a powdery *Oxyd* by the *Electrical Spark*.

Platina is Capable of being brought to a
state of fusion by burning it in Oxygen
Iron when burnt in Oxygen

produces a very pretty appearance
In the case of the Iron when burnt in Oxygen

Deposited the ~~gold~~ by Nitric Muricatic
Precipitated it by Sulphate of Iron.

Platina not acted on by Acids, excepting by
Nitric

Hydrogenetic Acid & Oxymuric Acid
In solution in Nitric Muricatic Acid may be precipitated
Muricatic of Ammonia

Muricatic of Platina may be changed
to

When Platina is dissolved in Oxymuric
acid it is Capable of being precipitated by

Solution of Muricatic of Ammonia
Muricatic of Ammonia

Solution containing Muricatic of Platina
Solution containing Muricatic of Platina

Obtain the Platina in its Metallic form
precipitated from Muricatic of

Ammonia. having obtained this salt you may
applying heat to this precipitate in a crucible

off & a drop of water will be left
which is the purest Platina

obtained quite Malleable.

Purchased in Bars at 15 ~~lbs~~ In Grains 3 or 4 Shillings 6d
The Color of Pure Platina is a sort of a whitish grey

very Ductile & Malleable Pressed may be into Plates &
drawn into Wire

Drawn into Wire in a Crucible & a drop of water
will be left which is the purest Platina

Dr. Marcet has an English Penny piece of it which is twice the
weight of a Silver one

~~Both the Potash and precipitate~~ (Plati-
but today does not ~~not~~

The Specific Gravity of Mall: Platinum is $21\frac{3}{10}$

Dr Wollaston succeeded in drawing a wire of $\frac{1}{10000}$ part
inch in Diameter, & which supported $61\frac{1}{3}$ without break
but Dr W succeeded in drawing one $\frac{1}{10000}$ part of an
(film is sp. Gravity only 11) The wire in the operation of
the wire is covered with silver which is dissolved in Nitric Acid
Solutions of Minerals of Platinum decomposed
by $\frac{1}{2}$ Mineral of Tin & other Metallic Salts
The Minerals of Tin deposits to it a bright red
or rather a yellow. The action of Caustic Alkali
at high Temp: is very obvious therefore can't be
in our cases of analysis

Platinum by fusion is capable of uniting
with many Metals, forming alloys
Gold & Lead & Silver.
It is not capable of combining with Sulphur

Platinum is not much used in Arts on
account of its high Price - it makes
exceedingly valuable crucibles on account of the
difficulty of fusion in Analytical Chemistry. Evapor-
dishes are made of it for small experiments

Platinum is used for $\frac{1}{4}$ touch holes of Guns
used to employed for that purpose
but is much softer than Platinum

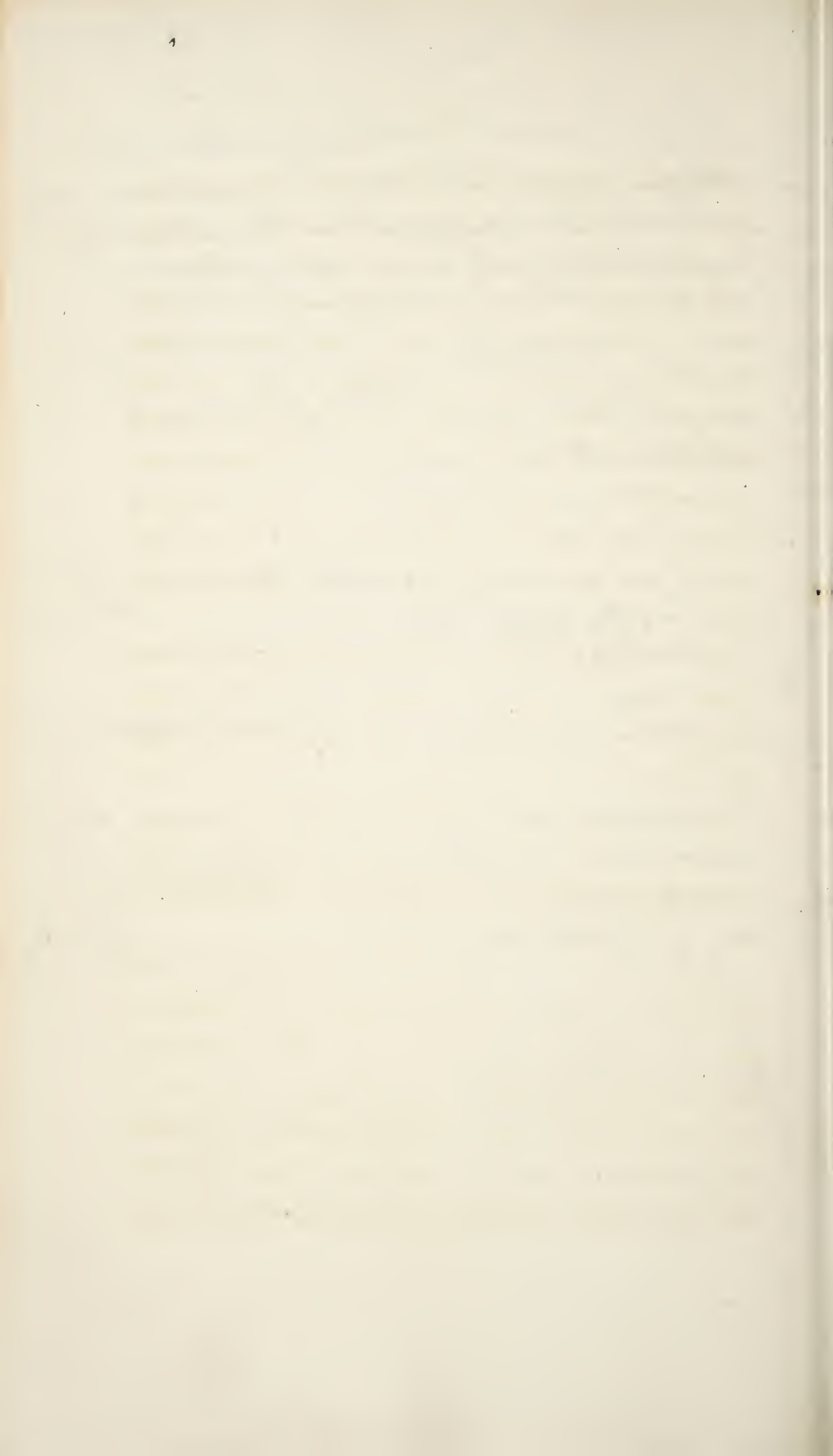
Mirrors for reflecting Telescopes are sometimes made
of Platinum

Platina vessels have been used lately
very much by Manufacturers - as
the preparation of Citric Acid.
Glass vessels but they are continually
rusting & are very expensive. The Platin
vessel on the other hand is used Bath
Metallic glazing which coats
thermore vessels, is made by $\frac{1}{2}$
of Platina - this glazing has a
pale white color

Platina may be fused by using the flame of a Spirit
lamp by means of Oxygen Gas. It is mixed in this way
with extreme facility. It was some time ago considered
impossible.

Platina at a white heat possesses the valuable property of
conducting

Platina is being capable of being annealed therefore
is useful in conducting molten metals. When
brought into the shape of a Spoon. Spoon tongs are
made also of Platina which are very useful



Soluble only in the *Nitro* and *Oxy-muriatic Acids*, to which it communicates at first a *yellow*, and afterwards a deep *reddish brown* colour.

Precipitable from its solutions in these by the *Alkalies* and several of their compounds—With *Muriate of Ammonia*, as with many of the others, it forms a *Triple Salt*, which on being submitted to an intense heat, and strongly hammered when in a state of incandescence, furnishes the pure *Metal*.—Precipitable also by many of the other metals and their solutions, more especially by *Tin*.

In its metallic state is not acted on, unless exposed to very high temperatures, by *Earths* or *Alkalies*, or any of their compound salts, except the *Nitrate* and *Oxy-muriate of Potash*, by which it may be superficially *oxydated* in the dry way.

Has no affinity with *Sulphur*; but, like *Gold*, is soluble in *Alkaline Sulphurets*—Combines readily with *Phosphorus*.

When pure, amalgamates intimately with *Quicksilver*, and unites by fusion, in different proportions, with most of the other *Metals*—most easily with *Zinc*—with *Copper* produces a golden-coloured, hard and durable compound, which is malleable, close-grained, and susceptible of a fine polish—remarkable for giving a

considerable degree of toughness to *Cast Iron*. With *steel* forms a compound incapable of being touched by the file.

Order of attraction, in the moist way, *Æther*, *Muriatic*, *Oxy-muriatic*, *Nitric*, *Sulphuric*, *Arsenic*, *Fluoric*, *Tartareous*, *Phosphoric*, *Sebacic*, *Oxalic*, *Citric*, *Formic*, *Lactic*, *Acetic*, and *Succinic Acids*; in the dry way, *Arsenic*—*Gold*—*Copper*—*Tin*—*Bismuth*—*Zinc*—*Antimony*—*Nickel*—*Cobalt*—*Manganese*—*Iron*—*Lead*—*Silver*—*Quicksilver*—and *Sulphuret of Potash*.

Use as yet, principally confined to *Chemical* and *Philosophical* purposes. Employed to make the touch-holes of guns—likely to be soon more extensively used,

Of Iridium, Osmium, Rhodium, and Palladium.

These four metals always found mixed or combined with crude platina. *Iridium* and *Osmium*, discovered in the black powder which remains undissolved after the action of nitro-muriatic acid on crude platina; and *Rhodium* and *Palladium* obtained from the solution itself.

Iridium first found in Mr Tennant's
- separated it from black powder by caustic alkali
- in mineral acids

properties are dissolved in all acids but
not in acid; insoluble...

Metal is obtained by heat alone not obtained

Metallic: Mass looks in a glossy mass.

as been found by the Galvanic Battery -

Osmium & ~~Rosicodendron~~ Iridium.

Osmium is obtained by heating the black powder
in silver crucible.

If a gentle heat by applying the Oxygen under
vacatilyze

A solution of Oxygen of Os. is colorless & does not
induce vegetable blues. If heat is applied the
Oxygen may be volatilized from it. it is volatile
at a lower Temp: than any other Oxygen of Metals.
Os. has never been got in Pure Metallic form
only in form of a Black powder.

Rhodium found in nickel Minerals. Soli
discovered by D^r Woll: precipitated by the
intermediation of Muriatic acid.

Correct Rhodium from Greek on account
of the rose color of its Salts.

Pure Rhodium is white

Specific Gravity only 11

Palladium may be got either from the same
solution by Precipitation of Mercury
Precipitating Palladium is from

100

100 grains of Platina only gives for Grain of
Palladium

In external appearance it resembles Pt.
it is very malleable more fusible
than Platina Minerals can do act for
it

I can burn with a sort of metals for
all alloys D. Woll: described a fine
age from

Solutions of *Iridium* varying in colour according to the degree of oxygination of this metal—How obtained pure. Soluble in acids, and especially in the muriatic. From its combination with this acid, may be obtained pure by heat alone.

Osmium, thus called from its peculiar smell. How obtained. Has never been procured in any other state than that of oxyd.—This oxyd very volatile and soluble in water. From this solution (which is colourless and transparent) the metal can be obtained in the form of a black powder. General properties of this substance.

Rhodium, thus called from the rose colour of its solutions. How separated from crude platina, and obtained in its reguline state. Its general properties.

Palladium, how obtained in its metallic state. Its general properties. Lately found in Brazil in grains of pure Palladium *mixed* with Platina, but not *alloyed* with it.

Of Gold.

Though less abundant, yet occurs more generally than most of the other Metals.

Found, either alloyed with a small proportion of Silver, Copper, Iron—*Native Gold*; or, combined with Sulphur, Antimony, Arsenic, Lead, Iron and Silver—*Grey Gold Ore*; or, with Bismuth and Sulphur—*White Gold Ore*—*Aurum Graphicum*.

Manner of collecting it from its ores.

When pure, of a *bright yellow* colour; *soft*; *inelastic*; *flexible*; *very tough*; *ductile* and *malleable* to an extraordinary degree; *not sonorous*.

Next to Platina the *heaviest* of the metals; its specific gravity when uncompressed being 19.30.

Scarcely *tarnishes* even by continued exposure to air or moisture.

Melts soon after becoming red hot, or at 32° Wedgw.— 5237° Farenht.—Whilst in fusion, of a *sea-green* colour.

Though unalterable in the common fire, may, by a more intense degree of heat, be *volatilized*, or imperfectly *oxydated*.

Crystallizes in cooling, into quadrangular pyramids.

If fused by the lowest degree of heat required for that purpose, becomes afterwards *brittle*—becomes also *hard* and *brittle* by compression; hence the necessity of *Annealing*.

Gold

considered above all other Metals -
account of its Ductility Malleability
and its not being acted on by $\frac{1}{2}$
atmosphere. -

exists in largest quantities in Mexico
anyway. - From the

$\frac{2}{3}$ quantity of in Ireland it is
and

found in flattened fragments of considerable
magnitude

has been discovered in an imperfect chrysolite
the more frequently found in Capillary State &
more frequently in the form of Dust.

When combined with some gold
you may Gold & Silver Leaf together and
put it into Nitric Acid you will find the
Silver acted upon but not the Gold.

of Gold may be extended 13000 Miles in length

an unbed with Silver may be separated by pouring
in Dilute Nitric acid

specific G: of Gold $19 \frac{3}{10}^{\text{th}}$

is any denser than that of water $\frac{1}{16}^{\text{th}}$ of an inch in
thickness will support

When Oure requires a White heat for its fusion
but when impure it will melt at a red heat.
It can't be Oxidated - only by the Galvanic Battery
Gold may be melted by means of a Spirit Lamp - which
is driven by a stream of Oxygen Gas - as Platinum may.
The Solvent of Gold is the Nitro Muriatic Acid
which is formed of Acs: Nit: 2 Parts. Acid: Muriatic
Part
Soluble in Oxymuriatic Acid

Solutions of Gold are of a rich Color

They are decomposed by γ Alkalies and the
Garners Compound called -

Fulminating Gold is formed by precipitating
a Powder, by Ammonia, from a solution of gold in
Nitro Muriatic Acid.
In the preparation of it the heat

The Powder should be placed on a card and held over
a lamp when it will explode violently. The explosion
is owing to the sudden formation of elastic Matter.
In precipitating γ Powder of Cassing - is formed
by putting Tin into a solution of Gold. Both the
Oxydes are precipitated in the form of a purple powder
also precipitated by the Sulphate of Iron in the
state of pure Gold - but it's not bright but can be
made so by means of Heat & Drying it. The
fewer never not precipitate the other metals out
Gold.

Gold has so little attraction for Oxygen that even the
alloy will reduce the Gold into its Metallic State

than cooling Gold it will change to alloy.

attraction of Gold for Oxygen is so weak that acids scarcely effect its solution.

Solutions of Gold are very corrosive they are precipitated by Alkalies

Solutions of Gold are decomposed when Gold is precipitated in its metallic form by some of the constituents.

Gold forms Alloys with most of the Metals

with Silver in

The Coin of this Country consists of 1 Part of Copper
to 11 of Gold

31 of Gold is worth intrinsically £3-12-0.

Gold unites with Ducksilver with great ease
forming an amalgam of a yellowish white
color this is much used in Gilding:

Gold has afforded another means of Gilding
as applied to enamel with size

Leaf is Gilded by

In the form of Gold leaf, is converted by *Electrical Explosions* into a *purple Oxyd*; and when ignited by the *Galvanic Fluid*, burns with *vivid white flame*, inclining to *blue*.

Like *Platina*, soluble only in the *Oxy* and *Nitro-muriatic Acids*, without effervescence; and in the solutions of *Alkaline Sulphurets*—Its acid solutions of a *yellow colour* and *caustic*, stain the skin *purple*, and, when evaporated, yield *deliquescent crystals*, which like the *oxyds* of this metal are decomposed by simple heat.

Gold separable from its solutions by *Æthers*, *Essential Oils*, *Phosphorus*, *Hydrogen Gas*, *burning Sulphur*, &c. May likewise be precipitated by all the *Alkalies*, and *Earths* in the form of a *yellow Oxyd*, which by exposure to light is partially decomposed, and becomes of a *purple colour*, and no longer soluble in acids.

The precipitate by *Ammonia* long known for the property of exploding with great violence when exposed to a moderate heat (*Aurum Fulminans*.)—This an *Ammoniacal Oxyd*, the explosive power of which depends on the double decomposition which takes place between the *Oxyd* and the *Ammonia*; hence the effects of exposing *Aurum Fulminans* to a gentle heat, or

of mixing it with *Oil* or *Sulphur*, or of heating it under strong *Compression*.

The precipitate of Gold by *Tin*—(*Purple Powder of Cassius*,) proved by the experiments of *Pelletier* to be an intimate mixture of *Oxyd* of *Tin* with imperfect *Oxyd* of *Gold*.

Gold distinguishable from *Platina*, by being precipitable from its solution by green *Sulphate* of *Iron*; but not by *Muriate* of *Ammonia*.

Its *Oxyds* unite by means of *Alkalies* with the *vitriifiable Earths*.

Has no affinity in its metallic state, for either *Sulphur* or *Carbon*; but unites with a small proportion of *Phosphorus*, and thereby becomes paler, and more fusible.

Combines with all the *Metals* in various proportions, acquiring different shades of colour and hardness, according to the quantity of alloy.

With *Copper* assumes a deeper colour and becomes more fusible, harder, and more elastic.

Amalgamates readily with *Quicksilver*; hence the art of water-gilding, and manner of collecting it from its ores, as before mentioned.

May be freed from admixture of imperfect metallic matter, by fusion with *Lead*, or *Bismuth* under free access of air; hence the process of *Cupellation*.

Instruction Value of pure Gold is £3-17-0
Answer -

leaf is gilded by putting the Part with
solution of Borax and heat afterwards
blinds

not on account of color that we used the
color of Gold but on account of its not being
lost upon by air & moisture







Most easily and effectually separated from *Silver* by diluted *Nitric Acid* : hence the process of *parting*.

Order of attraction in the moist way, *Æther*, *Muriatic*, *Oxy-muriatic*, *Nitric*, *Sulphuric*, *Arsenic*, *Fluoric*, *Tartareous*, *Phosphoric*, *Sebacic*, and *Prussic Acid*, *Potash*, *Ammonia* ; in the dry way, *Quicksilver*, *Copper*, *Silver*, *Lead*, *Bismuth*, *Tin*, *Antimony*, *Iron*, *Platina*, *Zinc*, *Nickel*, *Arsenic*, *Cobalt*, *Manganese*, *Sulphuret of Potash*.

Used extensively in the construction of *Uten-sils*, and different kinds of *Ornaments* ; in *Enamelling*, *Gilding*, *Dying*, *Soldering*. &c.

Of Silver.

Found, 1st. united to a small proportion of *Gold*, *Copper*, or *Iron*, *Native Silver* ;—2d. in alloy with *Antimony*, or with *Antimony*, *Arsenic* and *Iron*—*Antimoniated* and *Arsenical Silver Ores* ; 3d. mineralized by *Sulphur*—*Vitreous Silver Ore* ; 4th. combined, in the state of oxyd, with oxyd of *Antimony* and *Sulphur*—*Red Silver Ore* ; 5th. with *Lead*, *Sulphur*, *Antimony*, *Iron*, *Alumine* and *Silex*—*White Silver Ore* ;

6th. with Muriatic Acid, Oxyd of Iron, Alumine, and a little Sulphuric Acid—*Horn Silver Ore*.

Manner of extracting it from its ores.—Purified by *Cupellation* or reduction from *Muriate of Silver*.

Colour *white*—has neither *Smell* nor *Taste*—is less *ductile* and *tenaceous* than *Gold*; but *harder*, more *splendid* and more *opaque*—*Elasticity* between *Gold* and *Copper*—very *sonorous*—*Specific Gravity* from 10,253 to 11,091.

Tarnishes on exposure to air, and becomes occasionally encrusted with *Sulphuret of Silver*.

Melts below a white heat, or at 28 *Wedgw.* = 4717 *Farenh.* Is very *brilliant* when in fusion—and in cooling, *crystallizes* into quadrangular pyramids or octahedrons. By intense heat may, like *Gold*, be *volatilized* and partially *oxydated*.

By *electrical explosions* is converted into an *Oxyd* of a greenish grey colour—by the *Galvanic Fluid* burns with an *emerald-green* flame.

Its *Oxyds* decomposable by *Heat* alone.

Most readily *soluble* in *Nitric Acid*—the solution *bitter* and *corrosive*—stains the skin and other animal substances *black*.

Upon evaporation yields a crystallizable salt, susceptible of watery fusion (*Argenti Nitras*

The tubercular Silver ore is a Sulphuret of Silver

The red Silver Ore is a Compound of Antimony Silver Sulphur

The gray Silver ore is a Compound of lead Silver & Antimony

The Horn Silver is the form of a crust or a piece of Lead

Extraction of Silver from its ores is not difficult

It is of a Pure white color but tarnishes readily with Oxigen but from Sulphur

and is very malleable. The finest Silver leaf being made from the Point of an aitch in the hammer.

Ductility is greater but less than that of Gold and Iron

It is at 22 and 23 of Wedgewood's Pyrometer

It is fused may be made to crystallize in Tubes

It may be oxidized like Gold by Galvanic electricity

The Sulphure and to act on Silver requires
the assistance of Barley

Solutions of Argent. Nit. on exposure to light
deposits black particles which is Silver almost
in a Metallic State.

x Silver is evaporated to dryness it forms the
Luna Cornua.

The Silver may be obtained pure from the Luna
Cornua by heat.

The Luna Cornua appears to ^{consist of} 19 of Acid to
0.1 of Silver.

When the Luna Cornua is melted.

Nitrate And as scarcely any action on
Silver but microprecipitate it from solution.

Nitric Acid the most powerful solvent for Silver
but its necessity it should be Diluted with
water. This is remarkable, numerous bubbles
is which is owing to decomposition of the acid in contact
with the Silver. Nitrous Gas is evolved
oxide of Silver is formed. If the solution was
exposed to dryness.

Crystals of this salt when exposed to heat were
crystallized & hold a large portion of water of
crystallization
When its exposed to heat it fuses and is converted
into Argent, or Lunar Caustic.

Nitrate Acid readily unites with its Oxides
Silver & precipitates them. When the Muric of
Nitrate of Silver blackens on exposure to
the atmosphere,

Nitrate of Silver decomposed by other Metals
if a solution of this salt be applied to
plate of Copper it becomes coated with Silver
Arbor Diana is formed

Silver like Gold is precipitated from its solutions by.

Fulminating Silver is made by precipitating by Ag. Calces , and washing it with Ammonia in solution

It explodes on the slightest friction or heat, being applied to it. The explosion is owing to a sudden union of the Oxygen.

Take 40 gr of pure Silver. dissolved in 3j Anhydrous Nitric acid Ag. 3j Sp. Ues. 3j, ^{then gently} apply heat precipitates of a white powder, this is not inferior to the fulminating Silver. is detonated by Pressure Percussion in Contact with Ag. Sulphuric

Plating performed in a Mercury solution to Golding. The Silver is made into an amalgam with Quicksilver

Take of Black powder which is precipitated by Copper from solutions of Nitrate of Silver 4 parts Common salt & Sal Am. part: 10

Hyp: Axy: 3j mixed together & made into a Paste with water rub the surface of Copper with it Ag: 40 grs. and apply it

Another Method of Silvering
of the same Black Powder as above & Hyp: 10
Ues: 3j, Ammonia of part: 3j.

P. L.) Yields a *Metallic Precipitate* to many of the other *Metals*, as also to *Phosphorus*, burning *Charcoal*, burning *Sulphur*, &c.—and an *Oxyd* of silver to the *Fixed Alkalies* and *Earths*.—With *Ammonia* forms a triple compound still more remarkable for its fulminating property than that of *Gold* (*Argentum Fulminans.*) Manner of preparing this—and causes of failure.

Silver also soluble in Nitro-Sulphuric Acid (*Aqua Regina* of *Keir.*)

And in the common *Sulphuric Acid* by the assistance of a boiling heat.

Although slightly or not at all acted on by the other *Acids*, many combine readily with its *Oxyd*; hence the decomposition of *Nitrate of Silver* by *Muriatic* or *Oxy-muriatic Acid*, and their compounds, and the consequent formation of *Muriate of Silver* (*Luna Cornea.*) This remarkable for its easy *Fusibility*, *Insolubility* in water, and becoming *dark coloured* on exposure to *Light*, which produces a change of colour, equally striking, in *Chromate of Silver*.

Silver, in its *metallic state*, not combinable either with *Alkalies* or *Earths*, or with any of their saline compounds.

In the state of *Oxyd* it communicates a *yellowish olive* or *brown colour* to glass.

Unites in various proportions with most of the other *Metals*, and with all the other combustible *Bodies*, *Carbon*, *Azote* and *Hydrogen* excepted.—Loses its *Ductility* by combination with *Tin*, and with *Copper*, its usual alloy, becomes harder and more sonorous.—Forms a dark violet coloured mass with *Sulphur*.—With *Phosphorus* becomes more fusible and brittle.

Order of attraction in the moist way, *Muriatic*, *Sebacic*, *Oxalic*, *Sulphuric*, *Saccho-lactic*, *Phosphoric*, *Nitric*, *Arsenic*, *Fluoric*, *Tartareous*, *Citric*, *Formic*, *Lactic*, *Acetous*, *Succinic*, *Prussic*, and *Carbonic Acids*, *Ammonia*; in the dry way, *Lead*, *Copper*, *Quicksilver*, *Bismuth*, *Tin*, *Gold*, *Antimony*, *Iron*, *Manganese*, *Zinc*, *Arsenic*, *Nickel*, *Platina*, *Sulphuret of Potash*, *Sulphur*.

Used in the construction of various *Utensils*; in the composition of *Bell-metal*; in *Silvering*, *Enamelling*, *Soldering*, *Dying*, *Medicine*, &c.

Of Quicksilver.

Found chiefly either in a *Native State*; or alloyed with silver, *Native Amalgam*; or in union with muriatic and sulphuric acids, *Horn*

Black Powder 15 grains 3/4 Soda Ashes
Potash ^{carbonate} 3/4 5 Alums made into a
paste and spread over the metallic
surface you want to silver, but the
most substantial method of Silvering
is to mix Spirit Salting over the plate
of Borax, & lay on this a piece of lamel-
lar Silver & they are bound together that applied
Nitrate of Silver afford a very delicate test
or Mineral Acid — as also for Arsenic
and as an indelible Ink and by some
Darken of Hair also commonly used
a Cambric by Surgeons in this Country

Powder of Niter with Acid Sulph. P^{10} this compound
readily dissolves Silver

Silver can combine with Sulphur and also
with most of the Metals.

With Copper in certain proportions forms the
base coins of this Country.

From filings

Quicksilver

Sometimes found in Combination with other Metals

When pure it has a brilliant bright surface. the fluid opaque perfectly, reflect images. Sp. Gr. 13.6
39 below zero or

When exposed 71 Deg. below the freezing point it congeals, this experiment may be made by mixing Snow & Muscovy of Lime

Mr Walker however has shown that it may be congealed without Snow or ice by the action of cooling mixtures.

It may be congealed by the evaporation of Sulphur of Carbon or other a substance which is remarkable evaporable Dr. Marrett has succeeded with this liquor, filling with Quicksilver & wrapped it in cloths wetted with the fluid, & placed it in a chamber where the ^{sublimed} sulphur was used

Dr. Marrett applied the tube in which the frozen Quicksilver was contained, and applied it to his target when it produced a film of ice and found difficulty in detaching it. This is Dr. Marrett's explanation of this phenomenon.

Congealed Mercury undergoes much contraction directly opposite in that respect to Water $\frac{1}{23}$ of the whole bulk. it increases in size Cavendish expd the

Congealing Mercury gives an appearance of Crystallization.

Mercury boils at a Temp: of 650 or 660 its Vapor is visible like air, condenses again by cold: its Vapor is expanded like water by the operation of Caloric. Change of color Duckskin undergoes when exposed to air is owing to impurities being mixed with it. When by Distillation this change does not take place. Distillation produces a change of it. Makes it of a dark color. Most thought Oxygen was absorbed during this agitation. We may find that it was an Oxygen but Dr. Haer: does not say there is doubt about it. This is not thought in the case of Burgess's. The reason why it is not found that Oxygen is absorbed is because the Mercury is run from preparation spontaneously. Mercury mixed with Mercurous Chalk or fat are thought to be oxides as Empl: Hyd: Ung: Hyd: Oil Hyd: When Mercury is exposed to a boiling heat in a retort the Duckskin gradually loses its metallic luster it is converted into the Hyd: Oxyd: rubrum contains 8 parts in 100 of Oxygen by weight considered to be so. If the heat is continued the Oxygen is expelled again & the Duckskin is reduced again. Mercury acted upon more by heat, when only by heat Sulphur with which forms a Super Sulphate of Mercury the excess of Acid is driven off by heat it becomes Neutral Salt in this state it is soluble in water. If hot water be added to the Neutral Sulphate of Mercury it becomes yellow & a Sulphate of Mercury is formed.

owing to the Water attracting some of the acid from
the Salt. This yellow powder was called Turp
Mineral. Now Hyd. Sub fluoris or Idem bro.

Acid. Nitric: act powerfully on Mercury. If strong
the action is rapid. a White Crystalline is obtained.
Hyd. Nitrat. This is readily decomposed by
Ammonia by its stronger affinity for Nitric
acid.

Mercury; or mineralised by sulphur, *Native Cinnabar*.

Manner of obtaining it from its ores, and of ascertaining its purity. Freed most effectually from foreign admixture by cautious distillation.

When pure, of a *silver white* colour, *brilliant*, *fluid* at a common temperature, specific gravity 13.568.

Congeals at 39 below 0 *Farenh.* and then found to be malleable.

In its liquid form attracts moisture on exposure to the atmosphere, loses its splendor and contracts a grey pellicle. By agitation with access of air or trituration with mucilage or other tenaceous substances, is converted into a *greyish* or *black Oxyd*: hence the more common preparations of quicksilver, viz. *Pilul. Hydrarg. P. L. Unguent. Hydrarg. P. L. Emplast. Hydrarg. P. L. &c.*

Passes from its liquid state into that of vapour in vacuo even at a common temperature; but under atmospherical pressure, requires to be heated to 600 *Farenh.*

If continued under exposure to air at this temperature, is converted into a sparkling *red Oxyd*, containing a larger proportion of *Oxygen*

than that obtained by trituration, (*Hydrarg. oxyd. rubrum* P. L.) Of the chemical properties of this oxyd.—When exposed to a red heat in close vessels it is decomposed, yielding pure *Oxygen Gas*, and the original *Metal*.

Quicksilver acts either directly or indirectly on all the *Acids*.

Partially decomposes and combines with the *Sulphuric Acid* by the assistance of heat. Produces with it a white ponderous saline mass, which on the affusion of boiling water assumes a lemon yellow colour, (*Hydrarg. Vitriolat.* of the old Pharmac.) The change of colour thought to depend upon the abstraction of a portion of the undecomposed acid.

Dissolves in the *Nitric Acid* more or less readily, and with a more or less copious evolution of *Nitrous Gas*, according to the temperature and strength or dilution of the acid, *Nitrate of Quicksilver*.—This exposed to a low red heat, by a further and more complete decomposition of the acid, yields a red *Oxyd*, (*Hydrarg. Nitricooxyd.* P. L.) Analogous in all its properties to the *Hydrarg. Calcinat.* or common *Oxyd*.

Has no action on the common *muriatic acid*, unless previously oxydated; but with the *Oxygenised Acid* combines with great facility and

or Temp. of the water
forming the sub Sulphate of Mercury. This change is
due to the water attracting the Acid.

or as it is employed then is the Hypo-sulphate with a
very small quantity of water.

It is also known that the Acid should be
carefully kept & if kept in a glass bottle it is
not so good. Whether you employ dilute or pure Acid it is
not so good. It is best to keep the Acid in a glass bottle
and not in a metal one.

Hydrogen of Mercury is best kept in a glass bottle & the water
should be removed off & it is then known as a yellow
acid after this by continued heat it becomes
forming the sub Sulphate of Mercury and Hypo-sulphate of Mercury.

2. Mercurous oxide, the Acid is changed this
is very different from the Hypo: Oxyd: rubrum
which is not acted upon by Acid: Muriatic: unboiled in
State of Oxydation.

3. Oxymercuric soluble in 20 times its weight of water
this is very soluble in Alcohol.

(H) (NI) Hyd Oxygen

Liq Hyd Oxygen: Made by 6 viz Hyd
3i of Salution The Spirit promotes the solution

Hyd. Oxygen: Detected by very means
one of the best as viz. Dip a piece of Gold wire
a piece containing Hyd. Oxygen: Take a
piece of Zinc Wire & bring it into contact with
Gold. The Zinc

Mixture of Iron in solution is added a precipitate
will appear & a change of color to brownish
color.

When the Hyd. Oxygen is taken in large quantities
it produces Infl: of smaller intestines a very
don with sometimes even Salivation very quick
Dose $\frac{1}{10}$ of a grain to $\frac{1}{2}$.

Hyd. Oxy: as viz. Pumped Linct. 2 grains. Sulph. Ac
Mixture of Soda.

a quantity of Hyd. Oxym.: with carefully rubbed
with a weight of Lumps in the whole is
made of a greyish color. If this is heated &
sublimed a white powder is produced
which is Calomel. This is Sublimed 2 or 3
times before its use. If some Water
be added an Calomel by which it is deprived
a greater portion of its acid: it is Hyd. Oxym.
cum of Pure Ph: thought not to be likely to
be taken of Poison.

Ammonia also deprives it of its acid & makes
a dark color.

Antimony of Corrosive Subl. decomposed by
Alkalies. When the antimony is
decomposed by alkalis it is

oxide of Antimony PO. Subcarbon.

precipitating a solution of Hyd. Nit. by an
excess of Sulphureous, will form
precipitate analogous to Hyd. Submarine.

1000 of Mercuric in Sif of Acid. Nit: by that dissolved
into Cold Solution pour 3 1/2 of Alcohol by
Measure & apply that a fume arising the
fume falls down. This must be washed with
Cold Water afterwards dry it in a Water bath

Exploding force of Fulminating Mercury is produced

by the following process
Take 1000 of Mercury in Acid Nit: Sif & apply
by Measure the Solution which will
be 3 1/2 of Alcohol a fume arising
the fume falls down. This must be washed with
Cold Water afterwards dry it in a Water bath

without effervescence: hence the preparation of *Oxy-Muriate of Quicksilver*, (*Hydrarg. Oxy-Murias. P. L.*) and of *mild Muriate of Quicksilver*, (*Hydrarg. sub-Murias*; and *Hydrarg. oxyd cinereum, P. L.*)

Of the processes employed for obtaining these as well in the moist as in the dry way—new views of these compounds.

Of the combination of *Oxyd* of quicksilver with the *Acetic* and other *Acids*.

The acid solutions of this, like those of other *Metals*, decomposed by *Alkalies* and *Alkaline Sulphurets*, and by most of the *Earths*. The degree of previous *Oxydation* indicated by the colour of the resulting precipitates; those by *Potash* and *Soda* are of a yellow or reddish brown; those by *Ammonia* of a grey or white colour, (*Hydrarg. Præcipit. Alb. P. L.*) *Ammoniacal Muriate of Quicksilver*—FOURCROY.

Zinc, *Iron* and *Copper* precipitate *Quicksilver* from its solutions in its *metallic* form; a similar effect produced, though slowly, by *Phosphorus*.—*Alcohol*, added to the solution of *Nitrate of Quicksilver*, and assisted by heat, furnishes a precipitate, which when dried and heated, explodes with great violence, *Howard's Fulminating Mercury*.

Quicksilver amalgamates with most other *Metals*; very readily with *Gold*, *Silver*, *Lead*, *Tin*, *Zinc* and *Bismuth*; less easily with *Platina*, *Copper*, and *Arsenic*, and difficultly, if at all, with *Iron*.

Combines by different *Modes*, and, as was supposed, under different degrees of *Oxydation* with *Sulphur*. By fusion and subsequent trituration, or by trituration simply, into a ponderous black powder (*Hydrarg. cum Sulph.* of the old *Pharmac.*) *Æthiop's Mineral*; and by fusion and sublimation into a red striated mass (*Hydrarg. Sulphuret Rubr. P. L.*) *Vermillion*. *Artificial Cinnabar*. Which may also be prepared by double decomposition, from a mixture of *Muriate of Quicksilver* and *Sulphuret of Antimony*, *Cinnabar Antimonii*.

Preparations analogous to the former procurable, by agitating *Quicksilver* in solutions of *Alkaline Sulphurets*.

Order of attraction in the moist way, *Sebacic*, *Muriatic*, *Oxalic*, *Succinic*, *Arsenic*, *Phosphoric*, *Saccho-lactic*, *Tartareous*, *Citric*, *Nitric Fluoric*, *Acetous*, *Boracic*, and *Carbonic Acid*; in the dry way, *Gold*, *Silver*, *Platina*, *Lead*, *Tin*, *Zinc*, *Bismuth*, *Copper*, *Antimony*, *Arsenic*, *Iron*, *Alkaline Sulphurets*, *Sulphur*.

Goed With Duckstone is much
employed in Gelding

1 part of Sulph: only be mixed with 7 or 8 parts of
duckstone & sublimed. Vermilion is produced

duckstone is very important in arts &c

Preparat: of Mercury: as viz in Load: P

in its pure State.

1. Hyd Hyd Crut: Eluy: Hyd: in gross & diverse
perhaps can Oxidate

Hyd: Oxid: sublim

Hyd: Nitric Oxid:

Hyd: Oxymurias are Corrosive Sublimat

Hyd: Submurias are Calomel

Hyd: Oxid Ammon

Hyd: precipit: & Hous - which is Hyd Oxid
unprepared by chemistry:

Hyd: Sulph: sublim or Vermilion

Loses its lustre readily on exposure to *Air*, becoming first of a dull grey colour, and afterwards *whitish*.

Melts at 540 *Farenh.* and if cooled slowly crystallizes into quadrangular pyramids. Heated more intensely, it boils and emits fumes; if under exposure to air, it passes readily into a state of *Oxyd*, and assumes different colours, according to the degree of oxydation; hence *Grey Oxyd*, *Massicot*, *Minium* and *Litharge*. These and all the other oxyds of lead easily vitrified, *Glass of Lead*; and this easily decomposed, if heated, with the addition of *Charcoal*.

Lead in its metallic state little affected, either by the *Sulphuric* or *Muriatic Acid*, but dissolves readily in the diluted *Nitric Acid*, and forms with it a crystallizable salt.

Exposed to the vapour of the *Acetous Acid*, is converted into a laminated white oxyd, (*Flake White*, *Ceruse*, or *Plumbi Carbonas* of the last Pharm.), which, dissolved in a further portion of this acid, produces a crystallizable astringent salt, remarkable for its sweetness, (*Sugar of Lead*, *Plumbi Super-acetas*, P. L.) (*Liquor Plumbi Acetatis*, P. L.)

When highly oxydated, as by treatment of its red oxyd by *Nitric* or *Oxy-muriatic Acid*, forms

The Red Bay is by every heart given and the
Bay is the most famous in America and the town
which is now a city

and the Bay is now a city and the town
which is now a city and the town
which is now a city and the town

and the town is now a city and the town
which is now a city and the town
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and the town is now a city and the town
which is now a city and the town
which is now a city and the town

and the town is now a city and the town
which is now a city and the town
which is now a city and the town

and it is

Phosphate is only present by fermentation
in the soil. It is absent by Phosphate of Soda

If there is a dead in a pond the floating Anabaena
the white Camp is present. it is a Carbonate
dead

If Camp is washed in a pond - the true Salt
is formed the Phosphate is separated from it
it is a combination of the Camp with the Sodium

The big plant Anabaena ^{is not} is made by separating the
Anabaena from the water forming a compound

a *brown* precipitate, which inflames without detonation when strongly triturated with *Sulphur*.

Is precipitated from its solutions of a *white* colour by *Alkalies* and by *Earths*, and of a *dark brown* by *Alkaline Sulphurets* and *Sulph. Hydrog. Gas*. Its solutions also decomposed by the *Sulphuric, Muriatic* and *Phosphoric Acids*, and their compounds.

May be oxydated by deflagration with *Nitre*, and by fusion with the *Fixed Alkalies* becomes soluble in water.

Unites by fusion with most of the metals, viz. with *Platina, Gold, Silver, Copper, Tin, Bismuth, &c.* Amalgamates readily with *Quick-silver*, but refuses to combine with *Iron*.

Melts with *Sulphur* into a grey coloured brittle compound, less fusible than itself, *Artificial Galena*; with *Phosphorus* into one which is malleable, but more disposed to tarnish.

The *Oxyds* of *Lead* soluble in *Expressed Oils* and *Animal Fats*: hence the preparation of certain *Plasters, Varnishes* and *Paints*; are capable also of decomposing several of the compounded *Salts* (*Patent Yellow*); and remarkably promote the vitrification of earthy bodies and other metallic oxyds, as in *Glass-making*, and in the *Refinement of Gold and Silver*.

Order of attraction in the moist way, *Sulphuric, Sebacic, Saccho-lactic, Oxalic, Arsenic, Tartareous, Phosphoric, Muriatic, Nitric, Fluoric, Citric, Formic, Lactic, Acetous, Boracic, Prussic and Carbonic Acid, Potash*; in the dry way, *Gold, Silver, Copper, Quicksilver, Bismuth, Tin, Antimony, Platina, Arsenic, Zinc, Nickel, Iron, Alkaline Sulphurets, Sulphur.*

Employed in *Medicine*, and very extensively in the *Arts*, particularly in the construction of *Buildings*, and different *Utensils*; in the making of *Shot*, in *Statuary*, *Glass-making*, *Glazing*, *Painting*, *Varnishing*, *Refinement of Gold and Silver*, *Composition of Pewter*, and *Plumber's Solder*, &c. &c.

Of Copper.

Found 1st. *Native*; 2d. combined with oxygen and sometimes with iron, *Tile Ore, Pitch Copper Ore*; 3d. with carbonic acid, *Green and Azure Copper Ore*; 4th. with arsenic acid, *Arseniate of Copper*; 5th. with sulphuric acid, *Sulphate of Copper*; 6th. with muriatic acid, *Muriate of Copper*; 7th. with sulphur, *Vitreous Copper Ore*; 8th. with sulphur and iron, *Yellow*

The Masses which have been
by long sandy Opus

The Detection of Lead is very easy
blends, by some Preparation of Sulphur
the other fluid suspended in a
medium,

Copper

as known in most ancient times

ores of Copper are very common

Carbonate of Copper is also called Malachite

Copper sometimes is in Solution in form of the
Copper Mine

found in Wales in a kind of Tree which
contains a quantity of Copper, it was also observed
to the shoes of horses that fed on that Tree had
been with Copper

Combined with Arsenic forming Arsenic
Copper

When Copper is heated in Carbonate with other substances
it is reduced to metal by a further heat
than that of the carbonate.

Copper readily unites with the acids

By adding Ammonia to a solution of Copper
a white precipitate is formed

In order to combine Copper with Sulphur and Hydrogen
that is required in forming the Copper Sulphate of
the Electric Lamp. The chief application

Copper Ore; 9th. with arsenic and iron, *White Copper Ore*; and 10th. with lead, antimony, iron, sulphur, alumine, silex and silver, *Grey Copper Ore*.

Manner of *extracting* it from its ores, and of *purifying* it.

Colour, bright brownish red—*taste*, nauseous styptic—*odour*, when rubbed disagreeable—*hardness*, somewhat greater than that of *Gold* or *Silver*—*Malleability*, *Ductility* and *Tenacity* considerable—*Sp. Gr.* if soft 7.788, if compressed nearly 9—when hard, *elastic* and *sonorous*.

On exposure to *Air* loses its lustre and contracts a greenish rust: in the *open Fire* is converted into a dull brownish red or black *Oxyd*, which heated to redness with *Filings* of *Copper*, assumes an orange colour.

Fuses at 27° *Wedw.* = 4587 *Farenh.* and if heated more intensely is volatilized in fumes.—In cooling slowly *crystallizes* into quadrilateral pyramids.

Is more or less acted on by all the *Acids*.

Dissolves with the assistance of a boiling heat in concentrated *Sulphuric Acid*, and affords transparent oblong rhomboidal crystals of a deep blue colour, *Sulphate of Copper*, *Blue* or *Roman Vitriol* (*Cuprum Vitriolatum P. L.*)

Dissolves in diluted *Nitric Acid* with effervescence and the production of *Nitrous Gas*, forming a deep *blue* solution, and by evaporation a deliquescent salt, which detonates on being suddenly heated (*Nitrate of Copper.*) The precipitate obtained from this solution by adding *Chalk*, forms a beautiful but fugitive pigment, *Blue-verditer*.

Is in its metallic state slowly acted on by the *Muriatic Acid*, unless assisted by heat; the solution, of a grass green colour, on evaporation yields *cubical* crystals, whilst a solution of the orange coloured oxyd before mentioned, yields colourless *octahedrons*, which are not affected by *Ammonia*, unless after exposure to air.

Moistened with the *Acetous Acid*, under exposure to air, corrodes into a green saline oxyd, *Verdegris of Commerce* (*Ærugo P. L.*), which by additional acid dissolves, and yields a beautiful *dark green* transparent salt, *Distilled Verdegris*: both employed as pigments.

Easily attacked by the *Sebacic Acid* of rancid oil or fat, especially if before in any degree oxydated.

Most of the saline preparations of *Copper*, particularly the *Nitrate* and *Muriate*, soluble in *Alcohol*, which then burns with a green flame.

Sulphate of Copper has the property of depositing
acting violently on some of the Metals
when some of the Potent Salts is
added up in a piece of Tinplate. The tin
is accordingly dissolved to attract Oxygen,

its amount is not known unless added by
it

The Copper is very exposed to the fumes
of Benzoin. Benzoin is found,

the same of Ammonia or Sulphate of Copper
Parts Benzoin 1 Part Potash 1 Part
mercury 2 Parts When this mixture is
put in a crucible with a handle of glass



$\frac{2}{3}$ 3 Ounces of Copper filings

Hand and foot in an alloy of 1 Part of Copper
to 12 of Gold

With Hyd: it forms an alloy.

Is more or less acted on by all the *Alkalies*. With *Ammonia* forms a beautiful *blue* solution, which disappears on the exclusion and returns on the admission of *Air*. *Ammonia* the most delicate test of the presence of *Copper*, with the exception mentioned above.

Is precipitated of a fine grass green colour from its solution in *Sulphuric Acid*, by *Arseniate of Potash*, *Scheele's Green Pigment*. Of a *blue* colour from the *Nitric Acid* by *Chalk*, as before mentioned, and in a *metallic* state from all its solutions by *Zinc* and *Iron*, *Zement Copper*.

Detonates with melted *Nitre*, and by the assistance of heat decomposes *Muriate of Ammonia*.

Unites by fusion with many of the other *Metals*, forming very important compounds, as with *Platina* into one of great density and hardness; with *Gold*, *Standard Gold*; with *Silver*, *Standard Silver* and *Silver Solder*; with *Tin*, *Bronze* and *Bell-metal*; with *Arsenic*, *Tombac*; with *Zinc*, *Brass* and *Manheim Gold*, and with *Antimony* a violet coloured alloy.

Combines also with *Sulphur* and *Phosphorus* by fusion. Is tarnished by immersion in *hepatized Water*.

The *Oxyds* of *Copper*, impart a *greenish tinge* to *Glass*.

Order of attraction in the moist way, *Oxalic*, *Tartareous*, *Muriatic*, *Sulphuric*, *Saccho-lactic*, *Nitric*, *Sebacic*, *Arsenic*, *Phosphoric*, *Succinic*, *Fluoric*, *Citric*, *Formic*, *Lactic*, *Acetous*, *Boracic*, *Prussic* and *Carbonic Acids*, *Potash*, *Soda*, *Ammonia*, *Uctuous Oils*; in the dry way, *Gold*, *Silver*, *Arsenic*, *Iron*, *Manganese*, *Zinc*, *Antimony*, *Platina*, *Tin*, *Lead*, *Nickel*, *Bismuth*, *Cobalt*, *Quicksilver*, *Alkaline Sulphurets*, *Sulphur*, *Phosphorus*.

Use very extensive, being employed in the construction of *Buildings* and *Coppering* of *Ships*; in the formation of various *Culinary* or other *Utensils*, in *Bell* and *Cannon Founding*, in *Coinage*, in *Enamelling*, *Dying*, *Painting*, and *Medicine*.

Of Iron.

Of all metals the most frequently and abundantly met with, and generally in the following forms; 1st. *Native*.—Circumstances always attending the presence of *Native Iron*—*Native*

Iron

the susceptibility of Magnetism, by being attracted
to the iron in some of the rocks.

There is great variation but the attraction is not the same
in all circumstances.

Harvard.

It is to the Researches of an English Chemist, that we are
indebted for the knowledge of the properties of
this metal.

In Magnets we are sometimes surprised to find that

the Lodestone is always covered with a black crust
which is composed of Oxide.

The various parts of the magnet are connected together by a
Bridging Ridge of Iron & Nickel.

the same which fall have been found both the largest of
them nearly utilized

1000 at Madison in North America in the same way
by several persons, a correct account of them the
whole of them paper the whole. They weighed from 10
blasted in the 30 ball of Philadelphia. They are
generally very strongly facc obliquely which is
at they are penetrated with some considerable force
in some places

is supposed by some that they passed from some place
back in the past to some the same place. It is explained
that is merely hypothesis (This is Mr. Hancock's opinion)

the same is in greatest abundance in state of Oregon
where they are Red oxide of Iron or Hematite. on
account of its being used

in the Iron in Chazy State. Some count 2 north of the
Iron of the

some of them in Specimen of Hematite

of Iron is an oxide of Iron

oxide of Iron from Sweden which is the best

crystallized Iron Ore or compound of Iron

and Carbon and 8 Manganese

yellowish oxide of Iron

oxide of Iron or Sulphate of Iron which is very

common

To obtain Iron from the stone its necessary to heat
it to red heat & sulphur. The necessary one
is then put into a furnace with charcoal &
quantity of Iron is found with it

The more the charcoal is the more the Iron
The more the Iron the more the absorbing the
sulphur. & the more the sulphur the more the
Iron. & the more the Iron the more the
Iron. & the more the Iron the more the

Iron is more than the Iron

In order to convert the Iron into the
the metal & kept in a state of fusion & continued
the more the Iron the more the Iron
the more the Iron the more the Iron
the more the Iron the more the Iron
the more the Iron the more the Iron
the more the Iron the more the Iron

Iron proved to be of *Meteoric Origin*.—Description and chemical history of *Meteoric Stones*.—Origin of these stones discussed ; 2nd. in state of grey oxyd, *Grey Iron Ore* ; 3rd. united with carbonic acid, *Hæmatite* ; 4th. with carbonic acid, alumine, and often phosphoric acid, *Argillaceous Iron Ore* ; 5th. with carbonate of lime and oxyd of manganese, *Spathose Iron Ore* ; 6th. with sulphuric acid, *Native Sulphate of Iron* ; 7th. with chromic acid, alumine and silex, *Native Chromate of Iron* ; 8th. with sulphur, with arsenic ; or with both, *Iron Pyrites*, *Mispickel*, *Arsenical Pyrites*.

As obtained from its ores by the usual process of reduction, forms a fusible mass, of a blueish grey colour and coarse granular fracture, very hard and brittle. *Crude* or *Cast Iron*, which freed from its impurities by continued exposure to a strong heat and subsequent hammering and rolling, loses its brittleness, becomes of a lamellar or fibrous texture, and capable, when red hot, of having several portions welded by compression into a continuous mass, *Bar* or *forged Iron*.

This, by cementation or fusion with *charcoal*, acquires weight, becomes again fusible, brittle when cold, of a close granular texture, suscep-

tible of a high polish, very elastic and capable of taking on a great degree of hardness if heated and suddenly cooled, *Common and Cast Steel*.
 Operation of *Tempering*.

Bar Iron the purest. Colour of this *blueish grey*, when polished very *splendid*—has a slightly subacid taste, and when rubbed a sensible *odour*—*harder* than most other metals—more *tenacious* than any—considerably more *ductile* than *malleable*; Sp. Gr. from 7.6000 to 8.166.

Iron distinguished by its being attracted by the *Magnet*.

Susceptible of different degrees of *Oxydation*.
Black and Red Oxyd. Is speedily converted into a yellowish or reddish brown rust on exposure to air and moisture, (*Rubigo Ferri* of the former Pharm.), and still more speedily into a dark grey oxyd, by combustion in *Oxygen Gas*, digestion in warm *Water*, or the application of its vapour to it under ignition.

Dissolves more or less readily in all the *Acids*. Its solution generally accompanied with the evolution of *Hydrogen Gas*, and the resulting compounds possessing different properties according to its degree of *Oxydation*.

Requires the aid of heat to decompose *Concentrated Sulphuric Acid*, but dissolves readily in

Carbonaceous Iron

It is sometimes prepared by the following process

It is prepared by heating the ore in a furnace with charcoal for some time. The hot iron is then poured into a bucket.

Plunging red hot steel in cold water it increases hardness very much.

Temper steel you make it red hot then dipped in oil & taken out. The degree of hardness is judged off the color that appears if the blue color hardens it is as that of Springs of watches &c.

Properties
One of the most remarkable of iron is its magnetic property this applies principally to pure iron.

Iron is very susceptible of fusion heat of air contracts the carbon into pores quickly at a white heat iron is soft & extremely malleable. It acquires the malleable property of Wrought

iron by passing oxygen gas through it. It forms a black oxide of iron which by exposure to air absorbs more oxygen & becomes the red oxide of iron.

Oxygen may be obtained by heating iron filings in a glass jar.

Iron suffers by oxidation to air & rust. Common disease of iron is rust. It is a disease of carbon.

but when Iron reacts by application of heat
without acid Iron Hydro is formed.

In action of Sulphuric Acid on Iron evolved
Hydrogen



Precipitates spontaneously in the form of ochre from most of its acid solutions on exposure to *Air*. When thrown down from these by an *Alkali*, may be re-dissolved by the addition of a further portion of it, *Alkaline martial Tincture of Stahl*.

Combines with the *Fixed Alkalies* by fusion.

Deflagrates with *Nitrate of Potash*; and sublimes with *Muriate of Ammonia*, which it partially decomposes (*Ferrum Ammoniatum P. L.*)

When in the state of *Oxyd*, promotes the fusion of several of the *Earths*, and communicates different tinges to *Glass*, according to the degree of *Oxydation*.

Unites by fusion with all the other metals, except *Quicksilver*, *Lead*, and *Bismuth*. In combination with *Arsenic*, becomes brittle when heated, (*Red-short Iron*.)

Has of all metals the strongest attraction for *Sulphur*, with which, when heated, it unites very readily into a dark grey, brittle, and remarkably hard compound, *Artificial Pyrites*.

Is also capable of entering into combination with phosphorus, (*Cold-short Iron*), and with charcoal, *Steel* and *Plumbago*.

Order of attraction in the moist way, *Oxalic Acid*, *Tartareous*, *Sulphuric*, *Saccho-lactic Muri-*

verted into a *grey oxyd* (*Putty*;) by a continuance of the heat is changed into a perfectly *white Oxyd*, extremely difficult of fusion or reduction, *Basis of White Enamel*: a similar *Oxyd* obtained by exposing it when in fusion to the vapour of *Water*.

Broken when nearly or just congealed, exhibits an irregular columnar *Structure*, and by agitation while passing from the fluid to the solid state, may be reduced into very small grains (*Pulvis Stanni P. L.*)

Does not dissolve in the *Sulphuric Acid*, but by the assistance of heat decomposes it and becomes *oxydated*.

Rapidly decomposes the *Nitric Acid*, with a copious evolution of *Nitrous Gas*, and is converted into a *white Oxyd*: but in the considerably diluted acid, unaided by heat, dissolves and forms *Nitrate of Tin*, which burns with a thick white flame, and detonates with a heated crucible.

Dissolves readily in the *Muriatic Acid*; the solution yielding needle-form crystals which attract moisture.

Dissolves with still greater readiness in the *Oxy-muriatic Acid*, and in the *Nitro-muriatic Acid*, or *Aqua Regia*, yielding solutions remarkable for their property of forming a brilliant



and permanent scarlet precipitate with infusion of *Cochineal* and other analogous substances, (*Carmines*, and *Scarlet-dye*.) When united with *Oxy-muriatic Acid* in the *dry way*, by distillation with *Muriate of Quicksilver*, a volatile colourless liquor is obtained, which, on exposure to the air, emits copious dense white fumes, *Smoking Liquor of Libavius*.

Is slowly corroded by the continued application of the *Acetous Acid* in the form of vapour; and by long digestion, this acid dissolves a small proportion of it.

The solutions of *Caustic Fixed Alkalies*, assisted by heat, exert a solvent power both on *Tin* and its *Oxyds*, but most on the latter: *Caustic Ammonia* affects only the *Oxyds*.

Is little affected in the *dry way*, either by *Alkalies* or *Earths*, but decomposes the compounds of the former with the *Sulphuric Acid*. Detonates rapidly with *Nitre*, and decomposes *Muriate of Ammonia*.

Decomposes moistened *Nitrate of Copper* with an impetuosity productive of actual combustion.

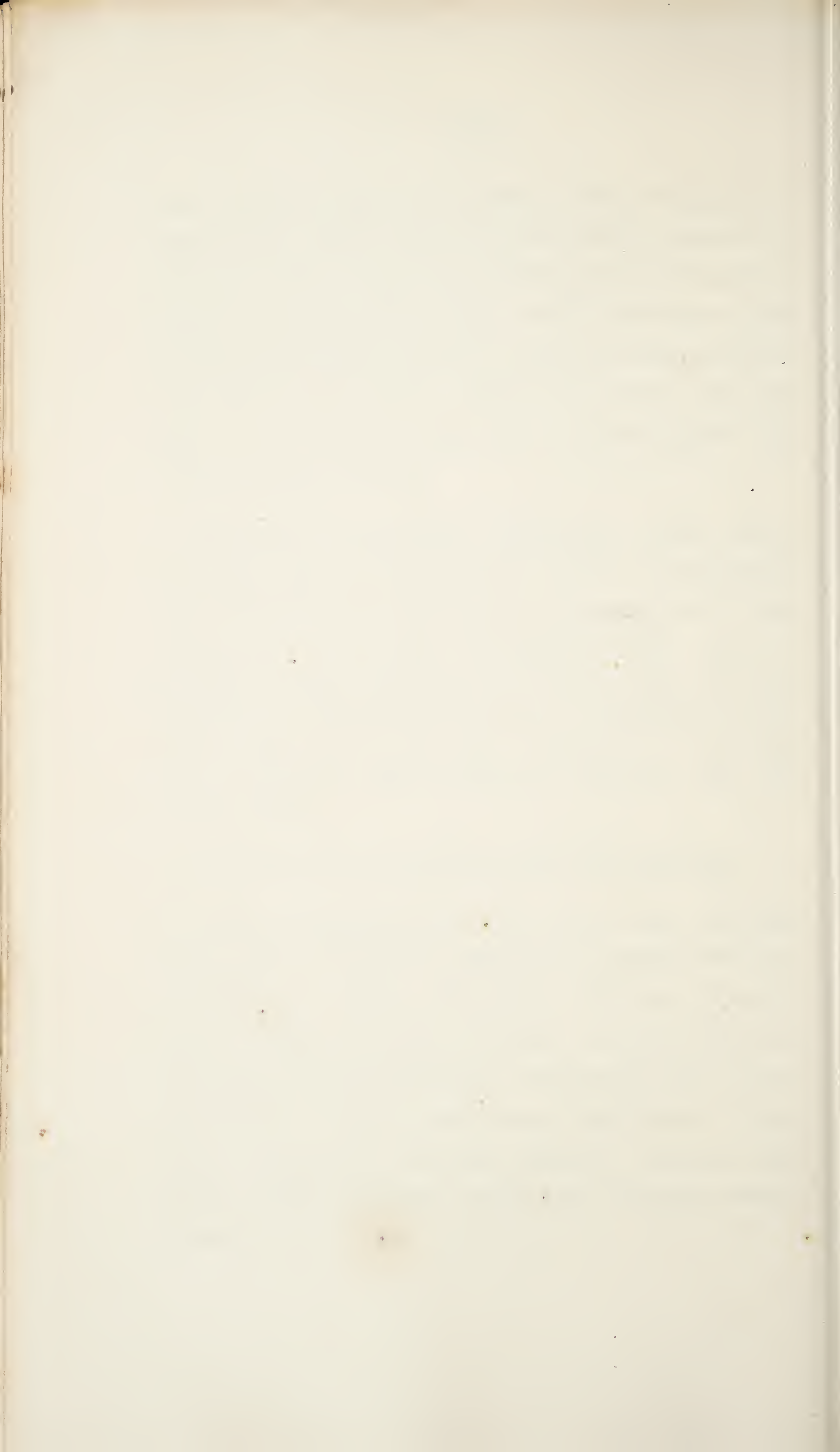
May be united more or less readily with all the other *Metals*, and in various proportions; occasioning in most of them an alteration in their *Malleability*, *Fusibility*, *Specific Gravity* and *Hard-*

ness : with *Copper*, according to the proportions, forms *Gun-Metal*, *Bronze*, *Bell-Metal*, *Speculum-Metal* ; with a small proportion of *Iron* becomes harder and more sonorous ; with half its weight of *Bismuth* forms *Tutenag* ; with an equal weight of *Lead*, *Soft Solder*. When free from *Arsenic* does not, in small quantity, much impair the malleability of *Gold*.

Melted with *Sulphur*, produces a dark grey, striated, brittle compound ; which, if prepared by sublimation, is of a light spongy texture and composed of golden coloured scales, used to imitate bronze, *Sulphuretted Oxyd of Tin*, *Aurum Musivum* ; which may also be prepared by heating gradually equal parts of *Sulphur* and the *White Oxyd*.

May also be combined with *Phosphorus*.

Order of attraction in the moist way, *Sebacic*, *Tartareous*, *Muriatic*, *Sulphuric*, *Oxalic*, *Arsenic*, *Phosphoric*, *Nitric*, *Succinic*, *Fluoric*, *Saccholarctic*, *Citric*, *Formic*, *Lactic*, *Acetic*, *Boracic*, *Prussic*, *Potash*, *Ammonia* ; in the dry way, *Zinc*, *Quicksilver*, *Copper*, *Antimony*, *Gold*, *Silver*, *Lead*, *Iron*, *Manganese*, *Nickel*, *Arsenic*, *Platina*, *Bismuth*, *Cobalt*, *Alkaline Sulphurets*, *Sulphur*, *Phosphorus*.





Employed in the composition of *Solder*, *Pewter*, *Bell and Gun-Metal*, *Printer's Types*, &c. in the silvering of *Mirrors*, construction of *Electrical Instruments*; in *Tinning*, *Enamelling*, *Dying*, *Medicine*, &c.

Of Bismuth.

Found 1st. *Native*; 2d. in union with *Oxygen*, *Native Oxyd*; 3d. with *Sulphur*, *Sulphuret of Bismuth*.

Manner of procuring it from its *Ores*.

Colour, *yellowish white*.—Specific gravity 9.822.—Texture *foliated*—not very brittle.

But little subject to tarnish. Melts at 460 *Farenh.* and in cooling crystallizes into cubes.

If kept in fusion with access of *Air* is converted at first into a blueish and afterwards into a brown *Oxyd*.—Heated to redness burns, with a small blue flame, and emits a yellowish smoke, condensable into an *Oxyd* of a similar colour, (*Flowers of Bismuth*,) which by increase of heat melts into a greenish *Glass*, analogous to glass of *Lead*.

Dissolves most readily in the *Nitric Acid*, and furnishes a salt, which detonates readily when exposed to sudden *Heat*; and which may be decomposed by the affusion of *Water*, yielding a white *Oxyd*, (*Magistery of Bismuth*). The other acids dissolve its oxyds, but do not act upon the metal.

Its precipitate by *Sulphuret of Ammonia* of a similar appearance to that which this produces in solutions of *Lead*.

Deflagrates both with *Nitre* and with *Oxymuriate of Potash*, and is converted into an *Oxyd*, which, like those already mentioned, facilitates the fusion of the *Earthy Bodies*, and also of the other *Metallic Oxyds*.

Combines with most of the other *metallic Substances*, rendering *Platina*, *Gold*, and *Silver* more brittle, and the metals in general more fusible.

Precipitates *Platina*, *Gold*, *Silver*, and *Quick-silver* from their acid solutions, but is itself precipitated by *Lead*.

Has very little affinity to *Phosphorus*.

Unites easily with *Sulphur* into a blueish-grey striated compound, which resembles the native *Sulphuret*, and may be decomposed by *Silver* and *Lead*.





Its *Oxyds*, like those of *Lead*, dissolve in *Unctuous Oils* by the assistance of heat.

Order of attraction in the moist way, *Oxalic Acid*, *Arsenic*, *Tartareous*, *Phosphoric*, *Sulphuric*, *Sebacic*, *Muriatic*, *Nitric*, *Fluoric*, *Saccho-lactic*, *Succinic*, *Citric*, *Formic*, *Lactic*, *Acetous*, *Prussic*, *Carbonic*, *Ammonia*; in the dry way, *Lead*, *Silver*, *Gold*, *Quicksilver*, *Antimony*, *Tin*, *Copper*, *Platina*, *Nickel*, *Iron*, *Zinc*, *Alkaline Sulphurets*, *Sulphur*.

Used in the composition of *Pewter*, *Solder*, and *Printer's Types*, in *Painting*, *Imitation of Silvering*, and occasionally in place of *Lead* in the refinement of *Gold* and *Silver*, and in *Assaying*. The *white Oxyd of Bismuth*, likely to prove an useful medicine.

Of Nickel.

Found 1st. alloyed with *Iron*, *Native Nickel*; 2^d. combined with *Oxygen*, *Native Oxyd*; 3^d. with *Iron*, *Arsenic*, *Cobalt* and *Copper*, *Kupfer Nickel*.

Extremely difficult to obtain it free from *iron*: accordingly varies somewhat in colour,

being occasionally *reddish, yellowish or greyish white*.—Specific gravity also varies from 7.88 to 9.333.—Texture *minutely granulated*.—Hard yet somewhat *malleable*. Is likewise *magnetic*.

Requires nearly the same heat as iron to fuse it; is very fixed in the fire, and difficult of oxydation by heat.

Deflagrated with *Nitre* yields a *greenish Oxyd*, which communicates a *hyacinthine tinge* to *Glass*.

All its solutions of a *green colour*: that in the *Nitric Acid* (in which it dissolves most readily) affords *green rhombic crystals*. Precipitated from its solutions by *lime water* and by the *alkalies*; *Ammonia* re-dissolves it, and produces a *blue solution*.

Forms brittle compounds with *Gold, Copper, Iron and Tin*; all the *Meteoric Stones* contain *Nickel* united with *Iron*; refuses to combine either with *Silver* or *Quicksilver*, and unites readily by fusion both with *Sulphur* and *Phosphorus*.

Order of attraction in the moist way, *Oxalic, Muriatic, Sulphuric, Tartareous, Nitric, Sebacic, Phosphoric, Fluoric, Saccho-lactic, Succinic, Citric, Formic, Lactic, Acetous, Arsenic, Boracic, Prussic, and Carbonic Acids, Ammonia*; in the





Arsenic

It is an oxide of Arsenic which we know under this name
Arsenical Pyrites obt^d from smelting the
In nature it is found in its Metallic State, then commonly
alloyed with Iron.

When Comb^d with Sulphur forms what is named Aspin
Realgar. Sometimes there are mixts the last is red & yellow
if this is treated with blowpipe it emits a Sulphur: smell.

Arseniate of Lime sometimes found but its a rare product

Pure arsenic brittle. Sp. G: 5.7

Passes into a State of Vapor without previous fusion & if
the process be performed without excess of air it condenses
at upper part of $\frac{2}{3}$ Vessel. Dr. Warron performs this in a
Glass Tube which is well cleaned: This succeeded well.

White oxide has an acrid taste & is corrosive. Soluble
in Water this passing in Oxymuriatic Acid & it
reddens the Inf. of Selenum & Comb: with $\frac{1}{2}$ Alkalies
Flowers of Arsenic.

Arsenious Acid is obtained by evaporation

dry way, *Iron, Cobalt, Arsenic, Copper, Gold Tin, Antimony, Platina, Bismuth, Lead, Silver, Zinc, Alkaline Sulphurets, Sulphur, Phosphorus.*

This Metal not yet applied to any use.

Of Arsenic.

Found 1st. alloyed by Iron, *Native Arsenic*; 2d. combined with Oxygen, *Native Oxyd*; 3d. with different proportions of Sulphur, *Realgar, Orpiment*; and 4th. with Sulphur, Iron, and sometimes Silver, *Mispickel*.

Colour, *steel-grey*—texture, *scaly*—very brittle, but not *hard*.—Specific gravity 5.763 to 831.—Much disposed to *tarnish*.

Volatile in close vessels at 356 of *Farenh.* and in condensing crystallizes in tetrahedrons.

If exposed to *Air* under the heat of ignition, burns with a blue flame, and is converted into a white *Oxyd, Common white Arsenic, Flowers of Arsenic, Arsenious Acid*.

This said to be soluble in about 80 times its weight of cold and 15 times its weight of boiling water. Taste *caustic*.—Less volatile than the *Regulus*.—Emits a garlic smell on the application of *Heat*.—*Vitrescible*.—Promotes the fusion

of *Earthy Bodies*. When melted with *Sulphur*, produces different shades of colour, according to the proportions and the degree of *Heat* or state of *oxydation*; hence artificial *Orpiment* and *Realgar*.—By treatment with the *Nitric Acid*, becomes more completely *oxygenated*, *Arsenic Acid*.—Decomposes *Nitre* in the dry way, producing a neutral crystallizable salt, *Arsenical Salt of Macquer*.—If distilled with unctuous matter, reassumes the form of *Regulus*.

In its *metallic* state, decomposes most of the mineral acids; the *nitric* most readily: if powdered and thrown into *Oxy-muriatic Acid Gas*, burns with a *white flame*.

Unites with most of the other *Metals*; rendering brittle those which are ductile, and in a remarkable manner increasing the fusibility of some, and diminishing that of others: thrown into *Oxy-muriatic Acid Gas*, burns with a *blue flame*.

To the red and yellowish metals communicates a silvery whiteness.

Soluble in *Alkaline Sulphurets*. Soluble also in *Uctuous Oils*, with which it forms a consistent mixture.

Combines readily with *Phosphorus*.

Most of the preparations of this metal in a high degree *noxious*; the detection of their pre-

By a very strong heat $\frac{1}{2}$ ars is decomposed again
becomes a white oxide

When melted with Sulphur in different proportions
forms different Chalzes.

Various salts are obtained as viz:

Hydrating Nitric with $\frac{1}{2}$ White oxide of arsenic
forms Arseniate of Arsenic:

Sulphate of Copper mixed with Arseniate of Potash
makes a fine Green color Scheele's Green

Known Capable of for alloys with most Metals destroys
their ductility & increases their fusibility.

Alloy of Arsenic & Platina is very fusible indeed

arsenic used in many of the arts

$\frac{2}{3}$ Manufacture of Glass it is employed also used in
dyeing.

Symptoms of Poison from Arsenic by Black
in a $\frac{1}{4}$ of Hour Sickness & Distress. & Heat in
Buccae involunt. Salivation a Bloody
fec^s from Mouth exs. pain & Nausea
Syncope. Cramps extreme weakness. &c.
Def: Inert: Infl: & Lamination crossers of
itself before Death. always the intestines
But be guarded in saying crossers of Stomach
are from arsenic as that takes place from
action of Gastric juice.

D. Marcet says it is mainly infl: of Mucous Coat
of the ^{Intestine} ~~Stomach~~ & infl: of Peritoneal Covering, from
irritations in Mucous Membrane of Intestine
Treatment Plentiful Drinks of Mucilage as Decar. &c.
De: Acid are the most efficacious steps to be taken
Water impregnated with Sulph: Hydrogen may be
usefully employed. if this can be borne give
Sulph: in Pills but better in Solution

Test: different ways proposed. by D. Marcet.
Take $\frac{1}{2}$ Suspended Sub: & put them into a Vessel,
the narrowest portion Section. Separated this
& mix it with a little Charcoal & Oil put it
in glass Tube & apply it to Heat. Then if it be
poisoned it will Chry. tallies at upper
part of the Vessel. a very small quantity may
be in this way detected. it produces a Metallic
ring on the Tube

a small quantity of the Metal may be mixed
with a little Charcoal & Oil & placed bet.
2 plates of glass & Red heat applied if ar-
senic present a white film will be produced

Arsenious acid is the most active poison
Blk. Oxyd: of Arsenic a deadly poison
Pure Metallic Arsenic is ineffective if alloyed
with another Metal if it is not it becomes
oxidized & therefore poisonous
In Solution much less Dangerous than when
taken in Substance. Sulphuretted Hydrogen comb'd
the water is a good antidote & Sulphur
arsenic is formed which is not very poisonous

the plate of Copper: If very thin plates of Copper
are employed a Spirit lamp will succeed but
if more employed the furnace.

we have not always a sufficiency of Matter to
obtain them readily. When they are made is
preferable. Act upon it in a State of Solution
the most delicate test in this Way is $\frac{1}{2}$ Nitrate
Silver. The Silver Comb: with $\frac{1}{2}$ Arsenic
forming a yellow precipitate a Little Alkali
Ammonia is necessary. a very small
quantity indeed may in this Way be detected
a large quantity of Ammonia would be
worrying... as the precipitate is soluble in
Solution of ammonia. Objections to this
there is a Mercaptic Salt we have have a
white precipitate discolouring the yellow one
oxide of Silver is produced. Phosphoric dis-
solving the same yellow precipitate with
oxide of Silver This has been thought an Objection
certainly is one. as it is not possible for
phosphoric acid to be preventing $\frac{1}{2}$ but this
objection may be removed as follows

Take 4 per cent of stannic form by Argenta
Nitrat: Mix it with Charcoal & supply
that a Metallic film will be found on it
by this means D.M. believes $\frac{1}{10}$ of a gr. can be
made apparent.

Sulphate of Copper forming Scheele's Green is also
a Test in a human way. if the solution of
Sulphate of Copper is at all acid, it requires
the addition of a little Ammonia.

Medically. Employed in Intermittent Fever. in
a shape of 4 Tasteless & Agreeable Drops now employed
in many dosages. The Dose is from $\frac{1}{12}$ to
of a 6 of 4 Hydri dechlorid and a solution of
Potash forming Resinate of Potash

Arg: stannic salt: it contains $\frac{1}{10}$ part
a Grain This Medicine is very useful
in Intermittents D.M. seems partial to
Objected to an account of its acceasing
Infl: Drathesis. When it disagrees causes
Pain in the Stomach Nausea Soreness & Inflammation
Swellings in the Stomach & Intest: as soon as these
Symptoms appear
Cobalt

Combined with Arsenic it is the grey Cobalt
One

sence therefore often of great moment. Various tests of Arsenic. Water impregnated with sulphuretted hydrogen, and solutions of Alkaline Sulphurets, recommended as antidotes. Emetics, purgatives, and copious dilution, the most effectual of these.

Order of attraction in the moist way, Muriatic Acid, Oxalic, Sulphuric, Nitric, Sebacic, Tartareous, Phosphoric, Fluoric, Saccho-lactic, Succinic, Citric, Formic, Lactic, Arsenic, Acetous, and Prussic, Ammonia, Unctuous Oils, Water : in the dry way, Nickel, Cobalt, Copper, Iron, Silver, Tin, Lead, Gold, Platina, Zinc, Antimony, Alkaline Sulphurets, Sulphur, Phosphorus.

Used principally in Glass-making, Painting, and Medicine. (*Arsenici Oxydum præparatum*; *Liquor Arsenicalis*, P. L.)

Of Cobalt.

Found 1st. alloyed with Arsenic, Grey Cobalt Ore ; 2d. united to Oxygen, Native Oxyd ; 3d. to Arsenic Acid, Cobalt Bloom ; 4th, to Sulphur, Sulphuret of Cobalt ; and 5th. to Arsenic, Iron, and Sulphur, White Cobalt Ore.

Colour, *light grey*. Specific gravity, from 7.7 to 8.53.—*Compact, brittle; hardness that of Tin.*

When very pure, malleable in a red heat.—Fuses at 130 *Wedgw.* and in cooling crystallizes in regular prisms.—Undergoes no change either on exposure to *Air* or *Water* at a common temperature; but, when heated in contact with air, yields at first a brownish oxyd, *Zaffre*, which on further exposure becomes blue, and on fusion with *Silex*, produces a blue glass, *Smalt, Powder-blue.*

Dissolves with more or less facility in several acids; most easily in the *Nitric Acid*.

With the *Muriatic* and *Oxy-muriatic Acids* forms compounds, the reddish solutions of which, as often as exposed to heat, change to a beautiful green, *Sympathetic Ink*.

Detonates feebly with *Nitre*.

Unites by fusion with most of the other *Metals*.

Combines difficultly if at all with *Sulphur*; more readily with *Alkaline Sulphurets*.

With *Phosphorus* forms a compound more fusible than itself.

Its *Oxyd* soluble in *Ammonia*.

Nature Oxide of Cobalt

It gives a blue color to any fusible substance

Not applied to any use except in a State of Oxide, therefore generally brought to that State

Cobalt in its pure State is of a grey color, but is slightly malleable, like Iron, and is attracted by γ Magnet.

Characters written with a solution of Cobalt are erasable when Cold, but are plainly seen when heat is applied. It loses its color again when it cools. Dr. Warrington thinks its action to the Ink is owing to its drawing off by γ application of heat. When dry it shows γ color.

Its use is to give a blue color to vitreous substances. When you suspect a substance to be Cobalt take a small piece of glass, Borax and the suspicious substance and if it is Cobalt it will turn the glass Borax Blue, which cannot be removed after the existence of Cobalt. A very small quantity will by this experiment be known.

Zinc

The Discovery credited to the Alchemists

Calamine an Oxide of Zinc.

Blind a Sulphuret of Zinc, but not a
pure Sulphuret.

Blind is often found in many parts of
the world but especially here

Calamine is of great use from which
Zinc is always obtained.

The surface is hard to the touch

specific Gravity $7 \frac{2}{10}$

Not Malleable (but little) or Ductile when
cold but at a certain temperature
is more so but at a high temperature
it is rendered again brittle and can be
powdered. forming granulated Zinc

Order of attraction in the moist way, *Oxalic Acid, Muriatic, Sulphuric, Tartareous, Nitric, Sebacic, Phosphoric, Fluoric, Saccho-lactic, Succinic, Citric, Formic, Lactic, Acetous, Arsenic, Boracic, Prussic, Carbonic, Ammonia*; in the dry way, *Iron, Nickel, Arsenic, Copper, Gold, Platina, Tin, Antimony, Zinc, Alkaline Sulphurets, Sulphur? Phosphorus.*

Used to colour Glass, in Glazing, Enamelling, Painting, Washing, &c.

Of Zinc.

Found 1st. combined with Oxygen, and frequently mixed with Oxyd of Iron, Silex and Alumine, *Calamine*; 2d. with Sulphur, Oxygen and other substances, *Sulphuret of Zinc, or Blend*; 3d. with Sulphuric Acid, *Sulphate of Zinc*; 4th. with Carbonic Acid, &c. *Carbonate of Zinc?*

Colour *blueish white*. Specific Gravity 7.190

Scarcely malleable, but can be rendered so by a particular process of annealing.

When heated is easily pulverised; and like iron is capable of Decomposing *Water*. Melts at about 700 of *Farenh.* Very easily volatilized.

In contact with air, burns soon after ignition, with a brilliant flame, and furnishes a white flocculent *Oxyd* (*Zinci Oxydum* P. L.) convertible by an increase of heat into a yellowish *Glass*.

Is readily acted on by *Acids*. During its solution in the *Sulphuric* and *Muriatic Acids*, *Hydrogen Gas* is evolved. Produces with the former Acid, a styptic crystallizable salt, (*Zinci Sulphas* P. L.) is also acted on by *Alkaline* solutions, and by *Water* previously impregnated with *Carbonic Acid Gas*.

May be precipitated from its solutions by *Earths* and *Alkalies*.

Decomposes, in the dry way, *Sulphate of Potash* and many other compounded salts. Detonates violently with *Nitre*. By simple trituration decomposes *Muriate of Ammonia*. Is also capable of precipitating *Alum* in the moist way.

Unites with all the other metallic substances except *Bismuth*; giving to *Copper* additional elasticity, durability and hardness.

In its *metallic form* cannot be combined with *Sulphur*; but when *oxydated* unites readily with it.

Order of attraction in the moist way, *Oxalic Acid*, *Sulphuric*, *Muriatic*, *Saccho-lactic*, *Nitric*,

When melted in contact with y Almas.
then it comes itself with a grey powder.

During its combustion the fumes of Iron
is formed.

From intense heat this oxide is vitrified
and if charcoal is present it is reduced.

There has scarcely any action on
water when cold but when heated
the Iron it decomposes it.

It is soluble in ² acids, when Sulphur
acid is poured on ^{Hydrogen} Iron, a gas will arise
which will burn, and when the flame
very small, it will produce a beautiful
kind of it be caused by a hollow
tube.

It detonates violently with Nitric

acid when combined with most of ²
metals

With Gold it changes its color to a
brake and destroy, its ductility

In a State of Sulphate it is used in
Medicine

Zinc.

Calamine is an Oxide of Zinc often combined with other substances.

When combined with Sulphur forms Blende or Sulph. of Zinc.

The color of Calamine varies: brown not a pure Sulph.
Comb^d generally from Lead arsenic &c
Blende & Calamine f^d in many parts of the World.

Zinc is generally obt^d from Calamine. It is powdered
Exposed to Heat with Charcoal. The Metallic matter
Condenses in $\frac{2}{3}$ Top of $\frac{1}{3}$ Vessel.

Zinc of a bluish white color.

Sp. G. $\frac{7}{10}$ ^{2th} Melted at 700° of Fahrenheit: kept that temperature
in closed Vessel it volatilizes without change.

If Zinc is heated it is Malable & Ductile. in high heat
may be broken with facility.

The flexibility by temperature in Zinc suggested
itself that it be made of it but this is now abandoned.
When Zinc is Melted in air it Oxidizes.

During the Combustion of Zinc $\frac{2}{3}$ Zinc. Axyd. is formed
which rises

Zinc has a power of decomposing Water when heated to redness -

Dilute Sulph: Acid acts upon Zinc powerfully & forms
Zinc Sulph: of Lead. Ph:

The Nitric & Muriatic acids act upon Zinc readily
with & last Hyd: Gas is formed?
Zinc readily deflagrates with Water.

Zinc in a state of Oxide combines with Sulphur
in forming amalgams with & Metals with Gold
& Silver which destroy its ductility.

With Copper it forms Brass - it is made by exposing
Alumina Charcoal & Copper to Heat & in this way
is readily formed

If a mixture of Zinc in a granulated state
mixed with Water be thrown into a red
hot crucible it burns splendidly.

Zinc is used in the construction of Galvanic
Cells.

Antimony.

a rare prod: in Nat State.

Other Metals occasionally found with it

Native Ant: of a yellowish white color. when heated with a Blow pipe gives out a peculiar smell.

The Sulphuret of Ant: is most abundant. Varieties of it
Red Antimonial Ore 2 in a Mass of Large crystals of
Red Antimony

Pure Antimony is obt^d by Melting the Sulph of Ant & then
when Separated is called Crude Antimony This
is purified by Heating it with Nitre.

Pure Ant: is of a bluish white color & when recently broken
it is destitute of Ductility Moderately Hard but yielding
to a Knife.

Sebacic, Tartareous, Phosphoric, Citric, Succinic, Fluoric, Arsenic, Formic, Lactic, Acetous, Boracic, Prussic, and Carbonic, Ammonia; in the dry way, *Copper, Antimony, Tin, Quicksilver, Silver, Gold, Cobalt, Arsenic, Platina, Bismuth, Lead, Nickel, Iron.*

Uses—in *Alloys*, particularly with *Copper* and *Tin*; externally and internally in *Medicine*.—A powerful agent in evolving *Galvanism*.

Of Antimony.

Found 1st. *Native*; 2d. alloyed with *Arsenic*, *Native Arsenical Antimony*; 3d. in union with *Oxygen*, *Native Oxyd*; 4th. with *Muriatic Acid*, *Muriate of Antimony*; 5th, with *Arsenic Acid* and *Sulphur*, *Red Antimonial Ore*; 6th. with *Sulphur* only, *Grey Antimonial Ore*; and 7th. with *Iron*, *Arsenic*, *Sulphur*, and sometimes *Silver*, *Plumose Antimonial Ore*.

Manner of extracting it from its ores, and the different processes for obtaining its *Regulus*.

Colour, *silvery-white*. Hardness, equal to *Zinc*. Texture, *laminated*. Very brittle. Sp. Gr. from 6.702 to 6.860. Simply loses its lustre

on exposure to the *Air*; and is not altered by *Water* unless exposed to it when red hot.

Melts soon after ignition; on cooling *crystallizes* into octohedrons; is *volatile* in close vessels. When heated in contact with *Air* is converted into a light white *Oxyd*, (formerly called *Argentine Flowers of Antimony*) which are soluble in water, and fusible into an *Hyacinthine Glass*; in close vessels by parting with different portions of oxygen, they acquire a *brown, orange, or yellow* colour.

Decomposes both with *Sulphuric* and *Nitric Acids*, the former with, the latter without, the assistance of heat. Requires long digestion for its solutions in the *Muriatic Acid*; but in the *Oxy-muriatic* dissolves with great facility.

Precipitable from the latter by *Zinc* or *Iron*, in the form of a *black Oxyd*, which when dried by a gentle heat *takes fire* spontaneously in the air, and is converted into *white Oxyd*.

Decomposes, in the dry way, most of the saline *Compounds* of the *Sulphuric Acid*. Detonates readily with *Nitre*; and decomposes *Muriate of Quicksilver*.

Combines with most other *Metals*, and renders them *brittle*. Said more than any other metal to diminish the *Magnetic* property of *Iron*.

the Metal suffers but little from Heat & Moisture but
in a White Heat or Oxidates & forms a Argentine
colour of Antimony. containing Dro 1/10 parts
of Oxygen.

Ant: disrobed by most Acids

Muriatic Acid acts slowly on Antimony but
acts powerfully on it when its Oxidates forming
Ant: Murias or Batter of Antimony. Made from
brown Oxide of Ant: & Common Salt with Sulph: Acid
Similar Comp^d may be formed by the connection
Oxygenated Acid with Antimony. it is highly
caustic when concentrated. If the Batter of Ant:
exposed to air it absorbs Moisture & becomes
acid.

Ant: Part: of Lond: Ph: Sulph: Ant: 2 Nitre,
Crown of Part: 2 chd. Sulph: 2 parts
The chd mixt with $\frac{1}{2}$ of Water & gently heated.

Chemical Affin chd: Part:

The Sulph: chd decomp: $\frac{1}{4}$ Nitre the Nitrous chd
set at liberty Oxyd: $\frac{1}{4}$ Ant: The exess of chd
comb. with Oxyd: of Ant: & forms Sulph: of
Ant: which is decomp: by $\frac{1}{2}$ Water
alone. by this process we have formed an
Oxyd: of Antimony wh^{ch} is capable of
uniting with $\frac{1}{2}$ Tartaric chd which
 $\frac{1}{2}$ Super Tartaric of Potash containing

Preparation now in use.

Ant. Axydum which is a Peroxyde.

Ant. Sulphur. Purif.

Pulv. Antimonial Sulphuret of Ant.
beated with Hart. horn shavings.

Ant. Tart.

Ant. Tart. Lignum 2 lbm & 1 lbm.

equal parts of Sulphur and of Antimony White disflav
forms $\frac{2}{7}$ Crocus of Antimony.

Calcined Ant: obtained from 1 part of $\frac{2}{7}$ Sulph. to 100 of
White. therefore very similar to the Above Crocus
Antimony

The Crocus has long been employed for $\frac{2}{7}$ diff
Antimonial preparations but now is in a
great measure exploded.

Unites with Sulphur in all proportions, and forms with it a grey striated compound, *Antimony* of the shops.

This, when exposed for a continuance to a low heat, yields a *Grey Oxid*, which by fusion is converted into a *yellowish Glass* (*Vitrum Antimonii* of the former Pharmac. ;) this levigated and mixed with melted wax forms the *Vitrum Antimonii Ceratum* of the late P. E.

Roasted with *Hartshorn* and afterwards ignited, yields a *white Powder* (*Pulvis Antimonialis* of the late P. L.—*Oxydum Antimonii cum Phosphate calcis* P. E.)—When deflagrated with *Nitre*, is more or less decomposed according to the degree of combustion (*Antimonium Calcinatum—Antimonium Ustum cum Nitro—Crocus Antimonii* of the former P. L. and P. E.)

Reduced to powder and boiled in a solution of *Potash*, deposits, on cooling, an *orange coloured precipitate* (*Kermes Mineralis* Ph. Suec.) ; and on the addition of the *Sulphuric Acid* to the solution whilst hot, a *brownish red precipitate*, (*Antimonii Sulphuretum præcipitatum* P. L.)
These *Hydro-sulphurated Oxyds of Antimony*.

A mixture of *Sulphuret of Antimony* and *Muriate of Quicksilver*, or of *Crocus of Antimony* and *common Salt*, with the addition of *Sulphuric Acid*,

produces, by distillation, a butyraceous compound, (*Antimonium Muriatum* of the late P.L.), which appears to be an *Oxy-muriate of Antimony*, and which on the affusion of common water, or by the addition of Potash, furnishes a *white Oxyd*, or rather *Sub-oxy-muriate*, (*Powder of Algaroth*.) The combination of this with acidulous Tartrite of Potash, forms a triple salt (*Antimonium Tartarisatum* of the old P.E.)

A similar preparation obtained from *Glass or Crocus of Antimony* (*Antimonium Tartarisatum*), which, like the former, is soluble in different kinds of Wine (*Vinum Antimonii tartarisatum* of the old Pharm.)

The preparations retained in the present Pharm. of London are *Antimonii Oxydum*, *Antimonii Sulphuretum præcipitatum*, *Antimonium tartarisatum*, *Liquor Antimonii tartarisati*. Nature of these preparations.

Order of attraction in the moist way, *Sebacic Acid*, *Muriatic*, *Oxalic*, *Sulphuric*, *Nitric*, *Tartareous*, *Saccho-lactic*, *Phosphoric*, *Citric*, *Succinic*, *Fluoric*, *Arsenic*, *Formic*, *Lactic*, *Acetous*, *Boracic*, *Prussic* and *Carbonic*; in the dry way, *Iron*, *Copper*, *Tin*, *Lead*, *Nickel*, *Silver*, *Bismuth*, *Zinc*, *Gold*, *Platina*, *Quicksilver*, *Arsenic*, *Cobalt*, *Alkaline Sulphurets*, *Sulphur*.

The Mercate of Antimony used to be
used as a Caustic but it is now
of little use from the Practice

Davy found the Mercate of Antimony contains
a mercuric Acid but is a compound
of Oxygenated Acid or Chlorine and
Antimony

If filings of Ant. be dropped into a Phos-
phoric Acid solution of Antimony it will be
oxidized and a Mercate of Antimony will be
formed

When Mercate of Ant. is concentrated by
evaporation, it becomes a powerful
caustic.

Both obtained by the action of Nitric on Antimony
simply by deflagration

The Ores is made by mixing --
equal parts of

The Ant. Calcinat. is part:

These three Oxides were formerly used as $\frac{1}{4}$
bases of the different preparation of Ant 7
but now there is a new Oxide introduced
by $\frac{1}{4}$ London College, which is considered
Superior.

Emetic Tart: a triple Combination of Antimony Potash
Tart. acid

The most important preparation of Antimony
is the Combination with $\frac{2}{4}$ Acid of
Tartar, made by boiling the
Super-tartrate of Potash -

with red brown Oxide of Antimony -
The Operation of very simple proportions,
Acids, Chlor. Chrys. of Tart. Chlor. Water Chlor. or
filtered, (Chlor. of fine S) and Chrys. Balling
it is the Antimon. Tartrazati

The Sol's Powder, appears to Phosphoric
Acid, Antimony and some combination
of these three things
1 part of Sulphur of Ant^m and 2 parts
of Cam Shaving Calcined together in broad
Pot.

Printing Ink is made by $\frac{1}{4}$ alloy of Ant: & Lead

If Sulphuret of Ant^y be boiled in a
solution of Potash ^{Diluted Water} and allowing it to
cool it forms y Kermes Minerales
or old Phannacopia.

Oxides of Ant^y are in some degree soluble
in urine it requires a tedious digestion
and after all the strength of different
preparations will be by no means
regular.

Ant^y forms alloys with other
metals, but the only useful one

or ^{or Naples} King's Yellow is Oxide of Antimony
and Oxide of Lead. ^(Antimonous) This is greatly
used in the preparation of Medicine
as a Sudorific. The antimon.
ant. is the one most commonly
used, and certainly one of the best
of its.

Ant: Calcanat. is a peroxide and al.
the oxide which is obtained.
consequently is not conforming with
the oxide. Antimony is a gentle
emetic.

Manganese

It is a metal which as a very strong action of Oxygen, that it is only found in a state of oxide

There are 2 different kinds of it.

1 in which it is combined with Oxygen & the other in which it is combined with Carbonic Acid

The Native Oxide found near 7 in England
The Carbonate called white found in England

The Purification of Mangan. is difficult on account of its impossibility. It has great attraction for Oxygen. It consists in exposing it to heat (intense with Carbon)

The silicious Manganese often combines with Quartz

Manganese neither Ductile or Malleable.

it is soluble in Nitric Sulphuric & Muriatic Acid: which solution may be decomposed by Alkalies & a White Precipitate on it.

The action of Acid. Nit. forms a Nitrous Gas & dilute

but the Sulph. acid. Muriatic and form Hydrogen Gas known to water decomposed.

Used in the composition of *Printer's Types*, and of *Nails for Coppering Ships*, in *Medicine*, &c.

Of Manganese.

Found 1st. *Native*; 2d, in union with *Oxygen*, *Native Oxyd*; and 3d. with *Oxygen*, *Silex*, *Iron*, and *Alumine*, *Siliceous Manganese*.—Said also to be contained in the ashes of most *Vegetables*.

Colour *dull* or *greyish white*. Texture, *granular*. Sp. Gr. from 6.850 to 7. Hardness next to *Iron*. *Very brittle*.

Next to *Platina* difficult to fuse; but *oxydates* more easily than any other metal: its *Oxyd* of different colours, *white*, *red* and *black*.

Soluble in the diluted *Sulphuric*, in the *Nitric*, *Muriatic* and several other acids.

In the state of *Oxyd*, occasions in the *Muriatic Acid* a striking change of properties, by imparting to it a portion of its *Oxygen*.

Its action on the *Alkalies* not yet ascertained.

When oxydated, decomposes both *Nitrate of Potash* and *Muriate of Ammonia*, in the dry way.

In this state unites by fusion with the *Earthy Bodies*, and when added to *Glass*, either renders it colourless or communicates a violet tinge, according to the degree of *Oxydation*.

Unites also, by fusion, with *Sulphur*, into a yellowish-green mass. By mixture with unctuous substances, sometimes occasions *Inflammation*.

Combines in the metallic state with most of the other *Metals*, rendering *Gold* and *Iron* more fusible, *Copper* less.

Order of attraction in the moist way, *Oxalic Acid*, *Citric*, *Phosphoric*, *Tartareous*, *Fluoric*, *Muriatic*, *Sulphuric*, *Nitric*, *Saccho-lactic*, *Succinic*, *Sebacic*, *Arsenic*, *Formic*, *Lactic*, *Acetous*, *Prussic*, and *Carbonic*; in the dry way, *Copper*, *Iron*, *Gold*, *Silver*, *Tin*, *Alkaline Sulphurets*.

Employed principally in *Glass-making* and *Bleaching*.

Of Tungsten, or Wolfram.

Found 1st. in an *acid* form in combination with *Lime*, *Tungstate of Lime*; and 2d. with *Oxyd of Manganese* and of *Iron*, *Silex*, and *Tin*, *Wolfram*.

Colour steel grey. Texture granular. Extremely hard. Sp. Gr. 17.6.

of Sulphuric Acid: is applied to Black
Manganese and heat applied Oxygen
gas is evolved

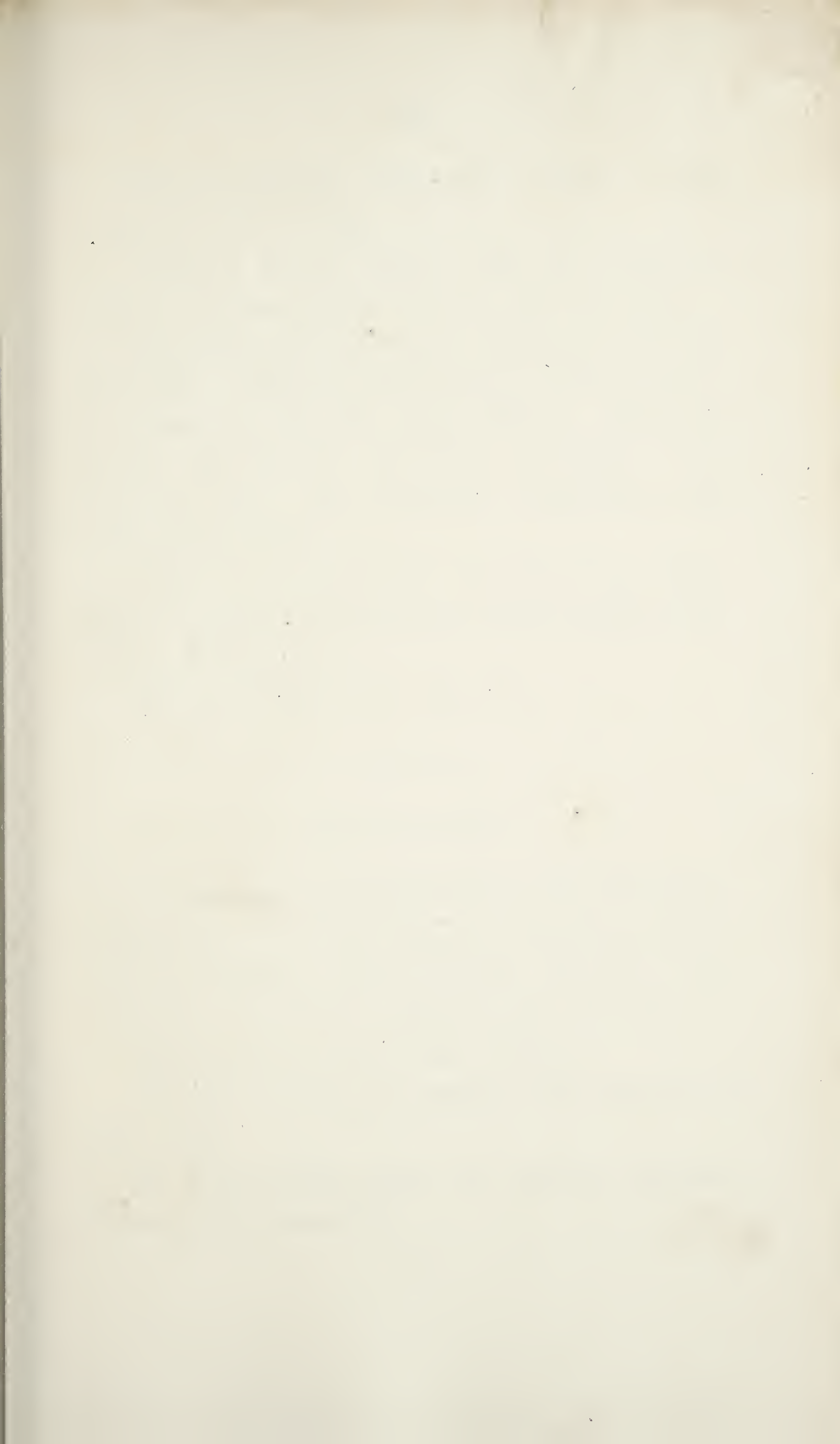
In the above Acid the Oxide of Manganese
is not dissolved unless a portion of
Hydrogen Gas be added

The first Alkali except an action ^{on} ~~of~~ ^{2nd} Black
Oxide of Manganese: they combine with
Heat and form a Compound, Soluble in
Water, and these Solutions can be
different colors by adding diff^t propor-
tions of Water ^{or diff^t temperatures} is called Water
and Pump Water used the pump Water
is not made rapidly by Distilled Water
is very imperfectly, Mineral Camellion
of Black Oxide and ^{2nd} Soda ^{or Potash} equal
parts, expose it to heat (red)

The same Compound formed by deflagrating
it with Black Oxide of Manganese in
Cucurbits not

^{Manganese}
Oxide, is used in the Manufacturing of
Lapis - as it is able to destroy the green
color which the Iron is apt to cause

The Oxymuratic Gas for Bleaching is made from
Black Oxide of Manganese.



Nearly *infusible*, requiring a heat at least equal to 170° *Wedg.*

Yields a yellow *Oxyd*, (*Acid of Tungsten*,) which in close vessels becomes *blue* or *black*.

Insoluble in the *Sulphuric*, *Nitric*, and *Muriatic Acids*. Slightly soluble in the *Oxy-muriatic*.

Combines with the other *Metals*. Does not lessen the ductility of *Silver* or *Copper*; but renders *Iron*, *Tin*, *Bismuth*, *Antimony*, and *Manganese* harder.

Order of attraction in the moist way, *Lime*, *Potash*, *Ammonia*; in the dry way, *Potash*, *Lime*, *Iron*, *Manganese*.

Not as yet applied to any use.

Of Uranium.

Found 1st. in combination with *Sulphur*, *Pechblende*; 2d. with *Oxygen*, (on the surface of the former,) *Yellow Oxyd*; and 3d. with *Carbonic Acid* and a little *Copper*, *Calcolite*.

Has been but imperfectly reduced.

Appears capable of uniting with several of the *Acids*.

Its *Oxyds* tinge glass of various colours, *brown*, *grey*, and *green*.

Of Molybdena.

Found in combination with Sulphur. *Molybdena*, formerly confounded with *Plumbago*.

Hitherto obtained only in agglutinated grains.

Colour, externally whitish yellow, internally grey. Sp. Gr. 7.5. *Brittle*.

Less fusible than either *Platina* or *Manganese*.

According to the experiments of *Mr. Hatchett*, is capable of combining with four different portions of *Oxygen*, producing a black, a blue, a green Oxyd, (*Molybdous Acid*,) and a yellow or white (*Molybdic Acid*.)

When combined with *Iron*, *Copper* and *Silver*, renders them friable.

In union with *Sulphur* regenerates *Sulphuret of Molybdena*.

Order of attraction unknown.

Not yet applied to any use.

Of Titanium.

Found combined with *Oxygen* in the *Red Schorl* of *Hungary*; and in *Manachanite*.

Of a colour somewhat resembling *Copper*. Appears to be *infusible*, but capable of being *volatilized*.

Difficultly *oxydated* by any of the acids.

With *Iron* forms a compound of a gold colour internally.

Of Tellurium.

Found in several of the *Ores of Gold*; and in considerable proportion.

Of a *whitish-lead*en colour; *metallic lustre*; *laminated* texture; *very brittle*; Sp. Gr. 6.115.

Of all metallic substances the most *fusible* except *Quicksilver*, like which it may be *sublimed* in brilliant globules.

Burns with a *blueish-green* flame. Soluble in the *Nitric*, *Muriatic*, and *Sulphuric Acids*; from which it may be precipitated by *Alkaline Sulphurets* in the form of a powder, in appearance much resembling *Kermes Mineral*.

Said to be the only metal, except *Gold*, *Platina* and *Antimony*, which is not precipitated from its solutions by *Prussian Alkali*.

Its *Oxyds* so rapidly reduced on heated *Charcoal* as to occasion actual *detonation*.

Of Chrome.

Found 1st. in an *Acid* form combined with Oxyd of Lead, *Red Lead of Siberia*; and in a similar acid form in the *pale red Ruby*; 2d. with Oxygen, in the *green Emerald of Peru*; 3d. in the *Meteoric Stones*.

May be obtained in a *concrete acid* state, and of a red colour, from the two first, by treatment with *Carbonate of Potash*; or in the form of a *green Oxyd* from the last by treatment with *Muriatic Acid*.

Has as yet been but imperfectly reduced.

Of a *greyish white* colour; *very hard*; *very brittle*; and *very difficult of fusion*.

Not soluble in the *Muriatic* or *Sulphuric Acids*; and difficultly in the *Nitric* and *Oxy-muriatic*.

Combined, in its *Acid* state with *Muriatic Acid*, is capable of dissolving *Gold*.

Of Columbium.

Discovered some years ago in an *Acid* state by *Mr. Hatchett*, in a dark grey mineral, sent with

some iron ore from *America*, and which appears to consist of more than three-fourths of this acid combined with iron. Its colour in this state is *white*; it *reddens* litmus paper; is insoluble in boiling *Nitric Acid*, but soluble in *Sulphuric Acid* when strongly heated, and also, when recently separated from Potash, in boiling *Muriatic Acid*.

It gives colourless solutions with *Acids* and *Alkalies*; is precipitated from its acid solutions by *Alkalies* in *white flocculi*, by *Prussiate of Potash* of an *olive green colour*, and by *Tincture of Galls* of a *deep orange*.

Is extremely difficult of reduction.

The supposed new metal called *Tantalum*, discovered by *Mr. Ekeberg* in the mineral called *Tantalite*, shewn by *Dr. Wollaston* to be identical with *Columbium*.

Of Cerium.

Lately discovered in a *copper mine* in *Sweden*. May be obtained in the state of *white* or *red Oxyd*. Both these oxyds soluble in *Sulphuric*

Acid. The white oxyd most soluble in the *Nitric*, and the red oxyd most soluble in the *Muriatic Acid*. The salts of *Cerium* have a *Saccharine* taste. Can hardly be said to have been obtained in the reguline state. Appears to be volatile at high temperatures.

ORGANISED BODIES.

Form two classes, *Vegetable* and *Animal*, consisting nearly of the same constituent principles or elements, but in different proportions.

The chemical affinities of the constituent principles of both, influenced by the *living principles*.

Carbon, Hydrogen and *Oxygen*, the *ultimate* constituent parts of *Vegetables*, as furnished by *Analysis*. *Air* and *Water* the principal sources of these. They also contain *Saline, Earthy* and *Metallic* matter, the latter in small quantity.

Manures. How they promote vegetation.

Circumstances in which *Vegetable* substances differ from *Animal*.

The Supply of Carbon is derived from 7 Alms.

The agency of Light is necessary for $\frac{1}{4}$ million of
vegetation

When the Sulphuric acid is used as Men-
struum in different drugs, Sassafras is usually the
residue. Sassafras for instance is useful
in the process of Pulverization.
If it be the carbonate of Sassafras is useful
on account of the Carbonic Acid Gas it
yields.

Vegetable Substances

These are formed naturally without any decomposition

Take the bark of the Quercus Suber graminea in the South of Europe like sand in its chemical nature is light Spongy Substances, inflammable affords a little Acetic acid by distillation it affords when acted on by γ Nitric Acid its yields the Suberic Acid.

Resins may be considered as bases of all other vegetables. naturally often possess of Taste and Color. The acetic to some essential Oil.

The Strong Acids act powerfully on the Acids. Sulphur: will completely alter it. Carbon is the Principal of

The Astringent quality of Suber graminea was at first thought to exist on the existence of Gallic Acid. Can contain in large quantities in Gall nut also in Cork. Quercus, and the mispropagation of γ Mucosa or Catechu. if you macerate the oak bark in Water the Tannin and Gallic Acid are dissolved by the Water of you put into this solution a little jelly of plainest precipitate takes place which is a compound of Gallic Acid Tannin and the Gallic Acid set at liberty

Vegetable Substances.

Their *proximate* parts, or immediate productions are, *Cork*, *Woody-fibre*, *Tan*, *Colouring Matter*, *Extract*, *Oils*, *Resins*, *Wax* and *Tallow*, *Camphor*, *Gum*, *Jelly*, *Gluten*, *Albumen*, *Caoutchouc*, *Starch*, *Sugar*, *Native Salts* and *Earth*.

Several of these already spoken of.

Cork,—the exterior part of the *Quercus Suber*, a substance *sui generis*. Yields *Suberic Acid* by treatment with the *Nitric Acid*.

Woody-fibre,—that which is left after the digestion of ligneous matter in *Water* and *Alcohol*.

Tan,—contained in all astringent vegetable substances, but most plentifully in *Catechu* and the *Gall Nut*, from the concentrated infusions of which it may be precipitated by *Carbonate of Potash* or by *Acids*. When dry is of a brown colour, brittle, of a very astringent taste, very soluble in *Water*, and still more so in *Alcohol*.—Forms an insoluble compound with *Glue*, and produces a deep blue precipitate when added to the solution of *Oxy-sulphate of Iron*. On distillation yields an *Acid Liquor*, a small proportion of *Empyreumatic Oil* and about *one-thirty-eighth* its quantity of *Charcoal*.

Colouring Matter,—that part of *Vegetable Substances* which is attracted by the fibres of *Silk, Linen, Wool, &c.* in the act of dying.—*Mordants*; their nature and use explained. Differs in its other properties according to the subject from which it is obtained.

Extract,—obtained from the watery infusion of *Saffron*, and many other plants, by evaporating it to dryness. Distinguishable from all other vegetable substances by being soluble both in *Water* and in *Alcohol*, but not in *Sulphuric Æther*.

Gum,—procured most plentifully from certain species of *Mimosa*. When pure, colourless, insipid, soluble in *Water* but not in *Alcohol* or *Oils*.—Not prone to decomposition by keeping.—Soluble in the *Vegetable Acids* without alteration, but decomposed by the *Mineral*. 100 parts said to consist of 65.38 *Oxygen*, 23.08 *Carbon*, and 11.54 *Hydrogen*.—An article of the *Materia Medica*; useful also in divers manufactures, particularly in *Calico Printing*.

Jelly,—obtained from the expressed juice of *Currants*, and many other fruits, in the form of *Coagulum*, by allowing the juice to remain for some time at rest, decanting off the thinner part, and washing the remainder in a small

an Infusion of Galls under form of Tannin
result

Tannin may be obtained by precipitating
a solution of by Lime water the
precipitate is Tannin mixed with
Lime which has to be got rid of
by Acid. Preparation:

Tannin soluble in water also slightly
in Alcohol

The most characteristic character of it is its
property of combining with Gelatin
thus the formation of Leather

This effect on Leather ascribed to an astringent
power exerted on the Leather in contact
with Gelatin. Mr. Sagar first took this view
of the Subject. The Tanning of Leather consists
in effecting a combination of Tannin and
Gelatin. The common way is to keep the skin
immersed in a solution of the Bark for
long periods.

The operation of Tanning may be very much
accelerated by putting the Leather into an
very strong Tannin rather than of oak
Bark. but Leather prepared in this way
is not found good, its only the outer
part of it that is affected. In Tanning
roan Leather, they only tan on skin by
the Sumach. The Russian Leather is tanned
by a very peculiar process, the color is given
by Siguum Campestre: it is almost ab-

Coloring Matter. Certain kinds of blue
or yellow Wood Madder. In the coloring
matter is given out to an aqueous
fluid. The coloring matter thus
formed has an affinity to attach
itself to linen.

But the coloring matter as a much
greater attraction for certain Mineral
Substances. Therefore a common
practice, to superimpose the cloth
dyed with some of these before the
color is applied. These are called Mordants
When a Mordant is used you can't wash
the color out which of course is an
advantage

Indigo a Beautiful blue Dye

It is not soluble in water but is in dilute
Sulphuric Acid: which is used as a
dye which acts at the same time as
a Mordant

The Vegetable Colours Called Lakes.

Carmine thus is also prepared by the same process, from Cochineal.

Extracts.. distinguished from all other vegetable Substances in being Soluble in Water

its general properties need not be said when Carmine is Soluble in Water and Alcohol (affinity for the Oxygen)

Gum is contained in many Plants but particularly in the Gumose species sometimes from the Bark sometimes from the Globules which are soluble in Water either Cold or Hot Spirit separates gum from a Solution of it. Gum differs from sugar only in containing only a smaller portion of Oxygen.

Vegetable Jelly. Somewhat resembles animal Gelatin in its translucent nature.

Gluten, contained abundantly in Wheat

When dried it is somewhat transparent & somewhat
gluey, it is on account of the Gluten
that it is so conveniently used in making
bread.

Albumen

Coagulable by heat, also in Acids

quantity of water.—In this state is nearly colourless, and of a tremulous consistence.—Soluble in hot water, but again coagulated in cooling.—When dried becomes transparent.—Combines readily with *Alkalies*, and by the *Nitric Acid* is converted into *Oxalic*.

Gluten,—the grey coloured tenacious and insipid substance which is left behind in washing the paste of *Wheaten Flour* in repeated portions of *Water*. In its *moist* state much disposed to putrify.—When dried resembles *Glue*.—In boiling water loses its tenacity. Soluble in all the acids, and in solutions of the *Alkalies* by the assistance of heat.—Insoluble in *Water*, *Alcohol*, *Æthers* and *Oils*.—Differs from most other *Vegetable* substances, and resembles *Animal* in containing *Azote*.

Albumen,—so called from its resemblance to white of egg.—Obtained by exposing to a boiling heat, the water employed in the preparation of *Starch* from flour, or the depurated expressed juice of *Scurvy-grass*, *Cresses*, *Cabbages*, and most other cruciform plants.—Separates at this temperature in the form of *Coagulated Flakes*.—Is also precipitated from its water solutions by *Alcohol* and *Acids*.—When dried resembles *Glue*, and is

then readily soluble in *Alkalies*.—In distillation yields *Carbonate of Ammonia*, and consequently contains *Azote*.

Caoutchouc,—the inspissated milky juice of certain trees, the *Hævea Caoutchouc*, *Iatropa Elastica*, and others growing chiefly in *South America*. When fresh is of an ash colour; without taste or smell.—Very elastic.—Sp. Gr. 0.933.—Not altered by *Air*.—Perfectly insoluble in *Water*.—Soluble in *rectified Petroleum*.—*Volatile Oils*, and in pure *Æther*, but not in *Alcohol*.—Is decomposed by the assistance of heat both by the *Sulphuric* and *Nitric Acids*, but not affected by *Alkalies*.—Melts readily when heated, yields *Ammonia* in distillation, and in the open fire burns with a white flame.

Starch,—the grey or white sediment deposited by *Whcaten Flour*, *Potatoes*, and various other vegetable substances, after diffusion in water.—Has scarcely any smell, and but little taste; though insoluble in *cold Water*, combines with it by *boiling* into a kind of *Mucilage*.—Altogether insoluble in *Alcohol*, and not readily acted on by *diluted Acids*; but soluble without decomposition by *Alkalies*, unless when they are concentrated and assisted by heat. By destructive distillation yields a large proportion of

Is a complete vegetable tissue

The grain is removed into a vessel from
rounds made into the form that produces
and moulds of clay in form of a little
bottle, the mould into the Juice which
is dipped into it. The stratum makes them
of the same process... very elastic... and a
property of its when cut is for
the cut edges to adhere firmly it is very
useful in Chemical experiments

Starch applied to a substance existing in
vegetables like Gum - produced by the
means for the separation of Glucose,
Totally insoluble in cold water but readily
converted into a Jelly by Boiling -
Pops Saloop & Lichen. Polakow. Polakow
in Wheat

It is considered the result of Saccharine
fermentation
It abounds in great abundance in many
vegetables. But the Sugar cane is the
thing which is planted for its selection
obtained by pressure and in the crude
state it undergoes a Process for
purifying it and forms the loaf
Sugar. Blood contains a large
portion of Saccharine Matter

Mr Howard has lately invented a new Method
of improvement.

a saturated solution of Symplic to use
for this purpose

Sugar may be considered a vegetable Symplic
Take 100 of Symplic 400 Water 6 of Acid Sulphuric
digest for 12 hours, filter and evaporate and
draw it alone, and it will become

Carbon. Supposed to be converted into *Sugar* in the process of *Malting*.

Sugar,—the produce of the *Arundo Saccharifera* or *Sugar Cane*, *Sugar Maple*, *Beet*, *Carrot*, *Grape*, and many other vegetables.—When purified, of a white colour, sweet, inodorous, not subject to alteration by exposure to *Air*; very soluble in water, and crystallizable; soluble also in *Alcohol*, but not in so large a proportion.—Readily decomposed both by the *Sulphuric* and *Nitric Acids*; in the weaker *Acids* merely soluble.—Forms with *Potash* a bitter astringent compound, insoluble in *Alcohol*; forms a similar one with *Lime*. On decomposition by heat yields *Water*, *Carbonic Acid*, *Carbonated Hydrogen Gas*, *Empyreumatic Oil*, *Acetous Acid*, and *Charcoal*.—100 parts consist, according to *Lavoisier*, of 64 *Oxygen*, 28 *Carbon*, and 8 *Hydrogen*.

Animal Substances.

Yield in general, by analysis, more *Azote* than *Vegetables*.

Their principal constituent parts and productions are, *Fibrine*, *Albumen*, *Gelatine*, *Mu-*

cilage, Urea, Sugar, Oils, Resins, Sulphur, Phosphorus, Acids, Alkalies, Earths, Metals.

Those not already treated of are,

Fibrine,—commonly called coagulable lymph, is that which remains on washing the crassamentum of blood in successive portions of water, till it cease to give out any colour. Bears a strong resemblance to muscular fibre ; is white, tasteless, soft, ductile, elastic, insoluble either in *Water* or *Alcohol* ; in boiling water becomes hard and inelastic ; in its soft state much disposed to putrify ; by long digestion in water is converted into a particular species of tallow.—Soluble in most of the *Acids*, and in the stronger solutions of both the *fixed Alkalies*, with which, by the assistance of heat, it forms a viscid bitter soap. In distillation yields a large proportion of *Ammonia*.

Albumen,—contained in the serum of blood, and in various other animal substances ; most abundantly in white of egg. In its moist and fresh state is nearly without colour, taste or smell—viscid—readily soluble in cold water—distinguishable from other animal fluids by coagulating at a temperature of 165 *Farenh.*—*Acids* and *Alcohol* produce on it a similar effect. When coagulated is insoluble in water.—*Tan*

The presence of Azote in Animal Matter
constitutes the difference between Animal
and vegetable Compounds.

Hence it happens that Vegetable Matter is so
liable to go into various fermentations
and animal Matter into the putrefactive
process.

Fibrine. If blood be removed from bone
its homogeneous look is soon destroyed and
separates

of this coagulum, be enclosed in a linen bag and
exposed to a stream of water, it will become colorless.

The Crasamentum is perfectly red.

The substance left after washing was formerly
called Glutin but now it is called Fibrin.

When the blood is replaced the Crasamentum
is covered with a coat of coagulum
lymph. The cause of this separation is unknown.
The Fibrine appears to form only 3 parts in 100.

When Dried it is a solid white substance
tasteless and without smell insoluble in
water and Alcohol when moistened and
exposed to Putrefy. Whites with $\frac{1}{2}$ Alkali
and forms a kind of soap.

The liquid separated from the blood by spontaneous
decomposition is commonly called Serum
but in chemical language it is called

Albumen, at a heat of 105 the fluid will be-
come opaque and have a look like the white
of an egg.

Albumen is soluble in water
and can be separated by Coagulation -
which is another striking character of Albumen.
Alcohol and Acids and several Metallic
Salts will coagulate the Albumen.

A solution of Hydr. Iod. is by far the most delicate test
for Albumen. Dr. Boston was the first who pointed out

A solution of Tannin slowly thrown down from
Albumen a yellow precipitate.

Albumen is Soluble in Alkalies

A very considerable excess of Acid has the power of precipitating its of depositing its Acid. It evidently decomposes it: it not only is destroyed by according to Mr. Wale's experiments it exists in Lig^t Cartilage. -
its Specific Gravity is various in different Specimens of it but in the greater number of Specimens Dr. Murrel examined

If Albumen which has been coagulated by heat is cut or flung is seen to exude from it which is called serosity and which is not at all susceptible of Coagulation. - This was generally supposed to be Gelatin - but now it appears that it possesses none of the properties of Gelatin and that it resembles Mucus more than any other Substance

Consist of Muriate of Potash and Soda.

The whole of Saline contents of Serum of blood is 1 grain of Saline Matter to 100 parts of Serum
100 gr of these Salinoids: 72 gr of Muriate of Soda
or 20 gr of Sod. Subcarb: or 10 gr of Potash Sulphate
Phosphate of Lime Iron and Magnesia.
The Sulphate of Potash is not found unless it has undergone incineration

The Alkali found in State of Subacromia

The quantity of the animal Matter in human blood
which when reduced to a dry brittle mass,

The specific gravity of human does not
depend on variation in quantity of salts, it depends
entirely on the quantity of animal Matter on
the coloring Matter of the blood, and that part
which may be washed off has been called

2. Red Globules, consisting of Albumen united
with Iron in a state of subphosphate to this gives it red color

Boards says the red Particles contain a
greater portion of ^{Iron} than of other Constituents
of the blood, this Dr. Hare doubts.

Gelatin or Jelly can be considered as composed
produced by animal substances

chiefly obtained from Skin by Tridons Caustic
lighter when the solution is allowed to

cool it becomes translucent - and is
known as when dried and its remaining

part dried off it is called animal
the name of Gelatin. it is quite hard & brittle

this was long believed to exist in the blood Dr. Boastock showed satis-
factorily this is not the case but it appears to be the result of a chemical process

Gelatin is insoluble in Alcohol - but
mixtures with Water will partially dissolve it
a solution of Isinglass in gum produces a

very good Paste for making various vessels
but it will not bear heat. The Best test

for Gelatin is Tannin this is the best
Dr. Boastock says 1 Part of jelly in 5000 parts of Water may
be detected with facility.

added to its aqueous solution, forms with it a copious yellow coloured tenacious precipitate, which, when dried, is brittle, and not susceptible of putrefaction.—*Albumen* soluble in *Alkalies* and *Alkaline Earths*.—In distillation yields the same products as *Fibrine*, but with a less proportion of *Ammonia*. *Tests* of the presence of *Albumen*. The fluid called *Serosity*, essentially different from *Albumen*.

Gelatine,—or Jelly, obtained by boiling certain animal substances, particularly skin, in common water, evaporating the decoction to the necessary degree of consistence, and allowing it to cool.—When pure, transparent and colourless. Soluble both in cold and hot water, very readily in the latter, even at a temperature of 90 *Farenh*. By drying becomes semi-transparent, hard, brittle, and of a vitreous fracture, *Glue*.—Is insoluble in *Alcohol*; *Alkalies* require the assistance of heat, but *Acids* dissolve it with great facility, even when diluted.—With *Tan* it forms, like *Albumen*, a yellowish coloured plastic compound, insoluble in water, and not susceptible of putrefaction; hence the theory of *Tanning*.—Heat decomposes it like other animal substances.

Mucus,—nature of this substance not yet well understood. Is distinguished by *negative* rather

*It has been often compared with Symplic
ing is with difficulty distinguished from other animal
substances.*

than by positive chemical properties. When exposed to the action of a voltaic pile, gives out Alkali at one pole, and a coagulated substance at the other.

Urea,—procured from fresh *Urine* in the form of small crystalline plates, by evaporating it to the consistence of syrup, digesting this when cold in *Alcohol*, distilling the solution so as to separate and collect the spirit, and allowing the residuum to crystallize by cooling. Thus obtained, its colour is *yellowish white*, smell *alliacious*, taste *strong and ammoniacal*, consistence *viscid*, extremely soluble in *Water*, somewhat less so in *Alcohol*, soluble also in the aqueous solutions of both the *Fixed Alkalies* and in *Marine Acid*.—By the *Sulphuric* and *Nitric Acids* it is decomposed, as it is likewise very easily by exposure to *heat*, even to that of boiling water. According to *Vauquelin*, 100 parts yield in distillation 92.02 *Carbonate of Ammonia*, 4.60 *Carbonated Hydrogen Gas*, and 3.22 *Charcoal*.

*Found to be
a new
Chemical*

Urea bears no resemblance to *Uric Acid*, a substance which constitutes the most common species of urinary calculus.—Conclusion of the Course.

THE END.

Perhaps there is no substance that has been
incorrectly defined as Mucus ... it is now
applied to a fluid secreted by a gland
set of glands to lubricate the parts
to which it is applied as the Urinary Mucus &c.
is in Alcohol soluble in water.

Mr Brander has inferred that Mucus and
Albumen are only modifications of same thing

mucus more and is required to coagulate
Albumen, there is required to neutralize
acid which it contains

The fluid of Notochord is an instance that they all
consist of Mucus dissolved in water. if you were
to evaporate a quantity of Urine, you will have
a solid substance, it can't be precipitated by
Tannin coagulable (not by Heat &c)

Urea may be obtained by evaporating a quantity
of Urine until reduced to a thick Symp. when
when all acid to cool Crystals will be formed

obtained in this method, it will crystallize

Urea is soluble in Alcohol and is obtained when this
is evaporated you will obtain Crystals of it

ly. Soluble in Alcohol and Water but the former
solution yields Crystals more readily than the water
when dissolved in water in diff^{er} proportions
it has the property of giving the various
kinds of different kinds of urine

If urine be saturated with Mucus of blood and
the solution crystallized, it will not form
its own regular crystals.

But there is no genuine test for it. It is
a compound.

D. Proust distinguishes two diff. kinds.
But he has not afforded route test
for it.

Calculi are forming in the kidneys particularly in
of Pelvis and Infundibulae

The symptoms indicating calculus in the kidney or in
the ureter going to the bladder, are very much alike.

Generally a considerable quantity of thick mucus
discharged with the urine.

The Presence of Stone in Bladder is principally indicated by a Pain at the end of $\frac{1}{2}$ Penis when the Urine is flowing a full stream & suddenly stops. There is also a continual inclination to make Water when the urine stops by the Stone stopping up the Passage & may be relieved by change of posture. All these Symptoms are increased by exercise.

Stone in $\frac{1}{2}$ Ureter.

When in the Prostrate Gland it is sometimes giving rise to difficulty of passing $\frac{1}{2}$ Urine but Calculi may exist for a long time without giving rise to any urgent symptoms.

The Symptoms of Calculus Concretions are in Women exactly similar to those of $\frac{1}{2}$ Male. We are indebted to Scheele for $\frac{1}{2}$ discovery of the fact of Calculi Lithic Acid - and he conceived that all of them must with was Lithic Acid. Fourquor Bonclay. & have brought the State of improvement in the knowledge that we now possess -

Those found in the Kidneys are generally from Lithic Calculi, on being cut thro they appear to be made up of various Lamellae with distinct Nucleus. But the Kidney has sometimes forming in it the Mulberry Calculus The Phosphate of Lime. Triple Phosphate.

Calculi in the Bladder are formed by various means, sometimes they consist of fine particles cemented together by Mucus - and of branches & ready getting into $\frac{1}{2}$ body will be liable to form a Nucleus of a Stone.

They are sometimes becoming of themselves loose.

The Shapes used Color also vary considerably
Some of them are smooth some rough
Some are white others Brown like the
color of a Mahogany;

Lactic Acid is disting by being Soluble in fixed
 Alkalies, ^{by applying gentle heat} and may be precipitated from
 this Solution by an Acid; if heated by
 plain Paper its entirely destroyed emitting
 - peculiar Smell. If Nitric Acid is drop

in it and heated to dryness. a beautiful
Crimson
Color is developed. which if you
add a little water ^{Crimson Solution} on ~~the~~ ^{formed} color
~~is imparted to it~~ when it is entirely

def. stone - Lythic Acid is very commonly met with
in the Urinary Bladder but not in a State
of Purity. This is most commonly met with in bad
Oxalate of Lime or Murexine Calculus
when it has it smells like Sperm.

Insoluble in Alkalies, difficulty in Acids
When urged by $\frac{1}{2}$ Blow Pipe it swells and
crackles - which is a test for it. The
Hydro Acid is burnt out by the heat and $\frac{1}{2}$
Lime remains in its pure state & will turn green.
If it is fully Purified and by $\frac{1}{2}$ application
of heat is soluble in Acids.

Phosphate of Lime is a Compact structure
Laminated and is radiated likewise when
may be seen by Close examination &
by that it remains unaltered excepting what
-ring it a little - Pure Phosphate of Lime

This is now asked upon by that subtle hand the
 The Fusible Calculus, when heated runs into a Globule
 Consists of an Ammoniacal: Magnesian Salt
 and Phosphate of Lime when these are separated
 they have peculiar properties. This Triple Salt
 when heat is applied emits an Ammoniacal Gas -

There is also feared, as I have written, that well as low has named Cystic Acids resembles very much the Hydrochloric Acid. Solutions of Acids injected into the Bladder has been tried but **A VIEW** not with much good effect. — **SIMPLE BODIES.*** Opium has also been dissolved in ~~any~~ Salutaria to allay the irritation of ~~the~~ **CLASS I.** *Bladder* —
 (Comprehending the Imponderable Agents, viz.)
 of course, right **CALORIC, or HEAT,** the Patient should be **LIGHT,** **ELECTRICITY.** *penicillin to injection being used.*

Ammonium the diff. kinds of Calcium with alkalies together. **CLASS II.**
 (Comprehending the Agents capable of uniting with Inflammable Bodies, and in most instances of effecting their Combustion, viz.)
these are **OXYGEN,**
OXY-MURIATIC ACID, or CHLORIC GAS.
CLASS III.

information (Comprehending the Simple Combustible Bodies, or Bodies capable of combining both with Oxygen, and with Chloric Gas, viz.)
HYDROGEN, or basis of Water.
NITROGEN, basis of Nitric Acid.
CARBON, basis of Carbonic Acid.
SULPHUR, basis of Sulphuric Acid.
PHOSPHORUS, basis of Phosphoric Acid.
BORACIUM, basis of Boracic Acid.
FLUORIUM, basis of Fluoric Acid.
POTASSIUM,
SODIUM, } **Metallic Bases of the Alkalies.**
AMMONIUM,†
CALCIUM, **YTTRIUM,**
MAGNIUM, **GLUCIUM,**
STRONTIUM, **ZIRCONIUM,** } **Metallic Bases of the Earths.**
SILICIUM, **BARIIUM,**
ALUMIUM,
PLATINA, **BISMUTH,** **MOLYBDENUM,**
GOLD, **NICKEL,** **TITANIUM,**
PALLADIUM, **ARSENIC,** **TELLURIUM,**
SILVER, **COBALT,** **CHROMIUM,**
MERCURY, **ZINC,** **COLUMBIUM,** } **Metals.**
LEAD, **ANTIMONY,** **CERIUM,**
COPPER, **MANGANESE,** **OSMIUM,**
IRON, **TUNGSTEN,** **IRIDIUM,**
TIN, **URANIUM,** **RHODIUM,**

* By denominating bodies *Simple*, it is not meant to express that they are so in reality, but only that we are not able, by any chemical means with which we are acquainted, to decompose them into simpler parts.
 † Ammonium is strongly suspected of being a compound; but as its nature has not yet been positively ascertained, it may still be classed amongst the simple bodies. The same may be said of sulphur and phosphorus.

ERRATA.

Page vii, Preface, line 8, instead of the word *occasionally*, read *usually*.

Page 37, line 3, from the bottom, dele the words, *soluble also in Spirit (Spiritus Ammoniae P.L.)*

Page 81, line 5, from the bottom, for *Wolfram*, read *Tungsten*.

Ibid. line 4, from the bottom, et passim, for *Chrome*, read *Chromium*; and after the word *Columbium*, add the words, *Cerium, Palladium, Osmium, Iridium, and Rhodium*.

Page 95, line 4, after the word *Quicksilver*, add, or *Calomel*.

Page 101, first line from the bottom, instead of *Cuprum Vitriol.*, read *Cupri Sulphas*, and add, *Triturated with Carbonat of Ammonia, forms the Cuprum Ammoniatum P.L.*

Page 108, lines 9 and 10, from the bottom, dele the words, *thrown into Muriatic Acid, burns with a blue flame*.

Page 119, line 8, from the bottom, for *præpartum*, read *præparatum*.

Page 125, at the end of the page, after the word *Charcoal*, add, *Mr. Hatchett's Artificial Tannin*.

